GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2004, 14:53:14; Search time 331.622 Seconds

(without alignments)

9184.983 Million cell updates/sec

Title: US-09-989-981A-9_COPY_3_104

Perfect score: 102

Sequence: 1 ctggtaggtgagatctctga.....aacaagctgtcctggaggcc 102

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

Database : EST:*

1: em_estba:*

2: em esthum:*

3: em estin:*

4: em estmu:*

5: em estov:*

6: em_estpl:*

7: em estro:*

8: em htc:*

9: gb est1:*

10: gb est2:*

11: gb htc:*

12: gb_est3:*

13: gb_est4:*

14: gb_est5:*

15: em_estfun:*

16: em_estom:*

17: em gss hum:*

18: em_gss_inv:*

19: em_gss_pln:*

20: em gss vrt:*

21: em_gss_fun:*

22: em_gss_mam:*

23: em gss mus:*

24: em gss_pro:*

25: em gss rod:*

26: em gss phg:*

27: em_gss_vrl:*

28: gb_gss1:*
29: gb gss2:*

ક

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

_	3 .		*				
Res			Query				B
]	No.	Score	Match	Length	DB	ID	Description
c	1	102	100.0	435	9	AI574075	AI574075 uj67h11.y
c	2	102	100.0	500	9	AI151811	AI151811 ui46c10.y
c	3	102	100.0	510	10	BB610072	BB610072 BB610072
c	4	102	100.0	511	9	AI157365	AI157365 ui45h01.y
С	5	102	100.0	583	13	BY705076	BY705076 BY705076
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C	7	102	100.0	3623	11	AK004871	AK004871 Mus muscu
C C	8	92.4	90.6	303	10	BB870338	BB870338 BB870338
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C	22	30.4	29.8	986	9	AV254401	AV254401 AV254401
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C	25	29.6	29.0	444	10	BF473385	BF473385 WHE0923_H
С	26	29.6	29.0	483	13	BQ467131	BQ467131 HS02L11r
С	27	29.6	29.0	518	14	CD912921	CD912921 G550.116E
C	28	29.6	29.0	564	12	BG606129	BG606129 WHE2960_H
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С	39	29	28.4	614	13	BU042469	BU042469 PP_LEa001
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С	43	28.8	28.2	342	9	AI117880	AI117880 uc41f02.r
C	44	28.8	28.2	398		AA177634	AA177634 mt32h12.r
C	45	28.8	28.2	416			BG550348 947039G04
_		-					

	46	28.8	28.2	510	13	BQ557757	ВQ557757 Н4048В01-
С	47	28.8	28.2	524	13	BX514645	BX514645 BX514645
С	48	28.8	28.2	536	13	BX520764	BX520764 BX520764
С	49	28.8	28.2	598	9	AI591944	AI591944 mt32h12.y
С	50	28.8	28.2	654	29	DR36H15S	AL987137 Danio rer

ALIGNMENTS

RESULT 1 AI574075/c AI574075 435 bp mRNA linear EST 29-MAR-1999 LOCUS uj67h11.yl Sugano mouse liver mlia Mus musculus cDNA clone DEFINITION IMAGE: 1925061 5', mRNA sequence. ACCESSION AI574075 AI574075.1 GI:4537449 VERSION KEYWORDS EST. SOURCE Mus musculus (house mouse) ORGANISM Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. REFERENCE (bases 1 to 435) AUTHORS Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R. TITLE The WashU-NCI Mouse EST Project 1999 Unpublished (1999) JOURNAL Contact: Marra M/WashU-NCI Mouse EST Project 1999 COMMENT Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@watson.wustl.edu This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. MGI:981353 Seq primer: custom primer used High quality sequence stop: 432. Location/Qualifiers **FEATURES** 1. .435 source /organism="Mus musculus" /mol type="mRNA" /strain="C57BL" /db xref="taxon:10090" /clone="IMAGE:1925061" /sex="female" /dev stage="adult" /lab host="DH10B" /clone lib="Sugano mouse liver mlia" /note="Organ: liver; Vector: pME18S-FL3; Site 1: DraIII (CACTGTGTG); Site 2: DraIII (CACCATGTG); 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTTTT]; double-stranded cDNA was ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested

and cloned into distinct DraIII sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should be used to isolate the cDNA insert. Size selection was performed to exclude fragments <1.5kb. Library constructed by Dr. Sumio Sugano (University of Tokyo Institute of Medical Science). Custom primers for sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end primer CGACCTGCAGCTCGAGCACA."

ORIGIN

100.0%; Score 102; DB 9; Length 435; Query Match Pred. No. 3.1e-22; Best Local Similarity 100.0%; 0; 0: Mismatches 0: Indels Gaps Matches 102; Conservative 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qу 166 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 107 Db 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102 Qу 106 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 65 Db RESULT 2 AI151811/c linear EST 30-SEP-1998 500 bp mRNA AI151811 LOCUS ui46c10.yl Sugano mouse embryo mewa Mus musculus cDNA clone DEFINITION IMAGE: 1885458 5', mRNA sequence. AI151811 ACCESSION AI151811.1 GI:3680280 VERSION KEYWORDS EST. Mus musculus (house mouse) SOURCE ORGANISM Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. (bases 1 to 500) REFERENCE Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., AUTHORS Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R. The WashU-HHMI Mouse EST Project TITLE Unpublished (1996) JOURNAL COMMENT Contact: Marra M/Mouse EST Project WashU-HHMI Mouse EST Project Washington University School of MedicineP 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@watson.wustl.edu This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. MGI:969782 Seq primer: custom primer used High quality sequence stop: 499. Location/Qualifiers FEATURES

1. .500

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                 with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTTTT];
                 double-stranded cDNA was ligated to a DraIII adaptor
                  [TGTTGGCCTACTGG], digested and cloned into distinct DraIII
                  sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site
                 CACCATGTG). XhoI should be used to isolate the cDNA
                 insert. Size selection was performed to exclude fragments
                  <1.5kb. Library constructed by Dr. Sumio Sugano
                  (University of Tokyo Institute of Medical Science).
                 Custom primers for sequencing: 5' end primer
                 CTTCTGCTCTAAAAGCTGCG and 3' end primer
                 CGACCTGCAGCTCGAGCACA."
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RESULT 3 BB610072/c

ORIGIN

Qv

Db

Query Match

EST 26-OCT-2001 LOCUS BB610072 510 bp mRNA linear BB610072 RIKEN full-length enriched, adult male liver Mus musculus DEFINITION

cDNA clone 1300007N20 5', mRNA sequence.

BB610072 ACCESSION

BB610072.1 GI:16451685 VERSION

KEYWORDS EST.

Mus musculus (house mouse) SOURCE

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 510)

AUTHORS Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T.,

Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J.,

Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,

Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,

Takeda, Y., Tanaka, T., Toya, T., Muramatsu, M. and Hayashizaki, Y.

RIKEN Mouse ESTs (Arakawa, T., et al. 2001) TITLE

```
Contact: Yoshihide Hayashizaki
COMMENT
           Laboratory for Genome Exploration Research Group, RIKEN Genomic
           Sciences Center (GSC), Yokohama Institute
           The Institute of Physical and Chemical Research (RIKEN)
           1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
           Tel: 81-45-503-9222
           Fax: 81-45-503-9216
           Email: genome-res@gsc.riken.go.jp,
           URL:http://genome.gsc.riken.go.jp/
           Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
           Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
           prepare full-length cDNA libraries for rapid discovery of new
           genes. Genome Res. . 10 (10), 1617-1630 (2000)
            waqi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
           Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T.,
           Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.
            and Hayashizaki, Y.
            RIKEN integrated sequence analysis (RISA) system--384-format
            sequencing pipeline with 384 multicapillary sequencer. Genome Res. .
            10 (11), 1757-1771 (2000)
            Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P.,
            Sugahara, Y. and Hayashizaki, Y.
            Computer-based methods for the mouse full-length cDNA
            encyclopedia: real-time sequence clustering for construction of a
            nonredundant cDNA library. Genome Res. . 11 (2), 281-289 (2001)
            Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I.,
            Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and
            Hayashizaki, Y.
            Computational Analysis of Full-Length Mouse cDNAs Compared with
            Human Genome Sequences. Mamm. Genome. 12, 673-677 (2001)
             Please visit our web site (http://genome.gsc.riken.go.jp) for
            further details.
             e mouse tissues.
                    Location/Qualifiers
FEATURES
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Qу
              228 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 169
Db
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Unpublished (2001)

JOURNAL

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DEFINITION
            ui45h01.yl Sugano mouse embryo mewa Mus musculus cDNA clone
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ACCESSION
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VERSION
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KEYWORDS
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 ORGANISM
           Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
               (bases 1 to 511)
            Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
 AUTHORS
            Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
            Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
            Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
            Waterston, R.
  TITLE
            The WashU-HHMI Mouse EST Project
  JOURNAL
            Unpublished (1996)
            Contact: Marra M/Mouse EST Project
COMMENT
            WashU-HHMI Mouse EST Project
            Washington University School of MedicineP
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:969717
            Seq primer: custom primer used
            High quality sequence stop: 480.
                     Location/Qualifiers
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                     Site 2: DraIII (CACCATGTG); 1st strand cDNA was primed
                     with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTTT];
                     double-stranded cDNA was ligated to a DraIII adaptor
                     [TGTTGGCCTACTGG], digested and cloned into distinct DraIII
                     sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site
                     CACCATGTG). XhoI should be used to isolate the cDNA
                     insert. Size selection was performed to exclude fragments
                     <1.5kb. Library constructed by Dr. Sumio Sugano
                     (University of Tokyo Institute of Medical Science).
                     Custom primers for sequencing: 5' end primer
```

CTTCTGCTCTAAAAGCTGCG and 3' end primer CGACCTGCAGCTCGAGCACA."

ORIGIN

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Query Match
                    100.0%; Score 102; DB 9; Length 511;
 Best Local Similarity
                    100.0%; Pred. No. 3.4e-22;
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 Matches 102; Conservative
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Qy
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ACCESSION BY705076

VERSION BY705076.1 GI:27116215

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 583)

AUTHORS Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C., Hume, D.A., Ouackenbush, J., Schriml, I., M., Kananin, A., Matsuda, H.

Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H., Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Brusic, V., Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglott, D.R., Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Pertea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S.,

Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M.,

Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E.S., Rogers, J., Birney, E. and Hayashizaki, Y.

```
Analysis of the mouse transcriptome based on functional annotation
 TITLE
            of 60,770 full-length cDNAs
            Nature 420, 563-573 (2002)
 JOURNAL
 MEDLINE
            22354683
            12466851
   PUBMED
            Contact: Yoshihide Hayashizaki
COMMENT
            Laboratory for Genome Exploration Research Group, RIKEN Genomic
            Sciences Center (GSC), Yokohama Institute
            The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            Tel: 81-45-503-9222
            Fax: 81-45-503-9216
            Email: genome-res@gsc.riken.go.jp,
            URL:http://genome.gsc.riken.go.jp/
            Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P.,
            Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F.,
            Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y.,
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            Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M.,
            Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y.
            Direct Submission
             Computational Analysis of Full-Length Mouse cDNAs Compared with
            Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
             Normalization and subtraction of cap-trapper-selected cDNAs to
            prepare full-length cDNA libraries for rapid discovery of new
            genes. Genome Res. 10 (10), 1617-1630 (2000)
             RIKEN integrated sequence analysis (RISA) system--384-format
            sequencing pipeline with 384 multicapillary sequencer. Genome Res.
            10 (11), 1757-1771 (2000)
             Computer-based methods for the mouse full-length cDNA
            encyclopedia: real-time sequence clustering for construction of a
            nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
             cDNA library was prepared and sequenced in Mouse Genome
            Encyclopedia Project of Genome Exploration Research Group in Riken
            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
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             Please visit our web site (http://genome.gsc.riken.go.jp) for
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REFERENCE
  AUTHORS
            Carninci, P. and Hayashizaki, Y.
            High-efficiency full-length cDNA cloning
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            Meth. Enzymol. 303, 19-44 (1999)
  JOURNAL
  MEDLINE
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            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
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            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
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            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
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            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
            Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M.,
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            The RIKEN Genome Exploration Research Group Phase II Team and the
            FANTOM Consortium.
            Functional annotation of a full-length mouse cDNA collection
  TITLE
  JOURNAL
            Nature 409, 685-690 (2001)
REFERENCE
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The FANTOM Consortium and the RIKEN Genome Exploration Research

AUTHORS

```
Analysis of the mouse transcriptome based on functional annotation
 TITLE
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
               (bases 1 to 2417)
REFERENCE
            Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
 AUTHORS
            Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
            Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,
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            Muramatsu, M. and Hayashizaki, Y.
  TITLE
            Direct Submission
  JOURNAL
            Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanaqawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
            URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
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            cDNA library was prepared and sequenced in Mouse Genome
COMMENT
            Encyclopedia Project of Genome Exploration Research Group in Riken
            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
            prepare mouse tissues.
            Please visit our web site for further details.
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Group Phase I & II Team.

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            Carninci, P. and Hayashizaki, Y.
            High-efficiency full-length cDNA cloning
  TITLE
            Meth. Enzymol. 303, 19-44 (1999)
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REFERENCE
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  AUTHORS
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
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            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
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            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
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            RIKEN integrated sequence analysis (RISA) system--384-format
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            sequencing pipeline with 384 multicapillary sequencer
            Genome Res. 10 (11), 1757-1771 (2000)
  JOURNAL
            20530913
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REFERENCE
            The RIKEN Genome Exploration Research Group Phase II Team and the
  AUTHORS
            FANTOM Consortium.
            Functional annotation of a full-length mouse cDNA collection
  TITLE
            Nature 409, 685-690 (2001)
  JOURNAL
REFERENCE
            5
            The FANTOM Consortium and the RIKEN Genome Exploration Research
  AUTHORS
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Analysis of the mouse transcriptome based on functional annotation

Group Phase I & II Team.

TITLE

of 60,770 full-length cDNAs Nature 420, 563-573 (2002) JOURNAL (bases 1 to 3623) REFERENCE Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H., **AUTHORS** Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M., Kasukawa, T., Kato, H., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C., Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Tanaka, T., Tejima, Y., Toya, T., Yamamura, T., Yasunishi, A., Yoshida, K., Yoshino, M., Muramatsu, M. and Hayashizaki, Y. TITLE Direct Submission Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of **JOURNAL** Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp, URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222, Fax:81-45-503-9216) Please visit our web site (http://genome.gsc.riken.go.jp/) for COMMENT further details. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. First strand cDNA was primed with a primer prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence[5' cleaved with XhoI and SstI. Cloning sites, 5' end: SstI; 3' end: XhoI. Host: SOLR. **FEATURES** Location/Qualifiers 1. .3623 source /organism="Mus musculus" /mol type="mRNA" /strain="C57BL/6J" /db xref="FANTOM DB:1300003C16" /db xref="MGI:1896857" /db xref="taxon:10090" /clone="1300003C16" /sex="male" /tissue type="liver" /clone lib="RIKEN full-length enriched mouse cDNA library" /dev stage="adult" 69. .2090 CDS /note="unnamed protein product; ATP-BINDING CASSETTE, SUB-FAMILY G, MEMBER 8 (STEROLIN-2) homolog [Mus musculus] (SWISSPROT|Q9DBMO, evidence: FASTY, 92%ID, 96.7%length, match=1796) putative"

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DEFINITION BB870338 RIKEN full-length enriched, adult male intestinal mucosa

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ACCESSION BB870338

VERSION BB870338.1 GI:17116548

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REFERENCE 1 (bases 1 to 303)

AUTHORS Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T.,

Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imotani, K., Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N.,

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Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A.,

Muramatsu, M. and Hayashizaki, Y.

TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)

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Unpublished (2001)
  JOURNAL
           Contact: Yoshihide Hayashizaki
COMMENT
           Laboratory for Genome Exploration Research Group, RIKEN Genomic
           Sciences Center(GSC), Yokohama Institute
           The Institute of Physical and Chemical Research (RIKEN)
           1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
           Tel: 81-45-503-9222
           Fax: 81-45-503-9216
           Email: qenome-res@gsc.riken.go.jp,
           URL: http://genome.gsc.riken.go.jp/
           Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
           Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
           prepare full-length cDNA libraries for rapid discovery of new
           genes. Genome Res. . 10 (10), 1617-1630 (2000)
            wagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
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           Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.
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            RIKEN integrated sequence analysis (RISA) system--384-format
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           10 (11), 1757-1771 (2000)
            Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P.,
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            Computer-based methods for the mouse full-length cDNA
           encyclopedia: real-time sequence clustering for construction of a
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            Please visit our web site (http://genome.gsc.riken.go.jp) for
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            Saurin, W. and Weissenbach, J.
            Estimate of human gene number provided by genome-wide analysis
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            Characterization and repeat analysis of the compact genome of the
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  JOURNAL
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            - Web : www.genoscope.cns.fr)
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            Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imotani, K.,
            Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M.,
            Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R.,
            Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N.,
            Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T.,
            Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
            Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A.,
            Muramatsu, M. and Hayashizaki, Y.
            RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al.
  TITLE
            2001)
  JOURNAL
            Unpublished (2001)
COMMENT
            Contact: Yoshihide Hayashizaki
            Laboratory for Genome Exploration Research Group, RIKEN Genomic
            Sciences Center(GSC), Yokohama Institute
            The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            Tel: 81-45-503-9222
            Fax: 81-45-503-9216
            Email: genome-res@gsc.riken.go.jp,
            URL:http://genome.gsc.riken.go.jp/
            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
             Normalization and subtraction of cap-trapper-selected cDNAs to
            prepare full-length cDNA libraries for rapid discovery of new
            genes. Genome Res. . 10 (10), 1617-1630 (2000)
             wagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
            Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T.,
            Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.
            and Hayashizaki, Y.
             RIKEN integrated sequence analysis (RISA) system--384-format
            sequencing pipeline with 384 multicapillary sequencer. Genome Res. .
            10 (11), 1757-1771 (2000)
             Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P.,
            Sugahara, Y. and Hayashizaki, Y.
```

```
encyclopedia: real-time sequence clustering for construction of a
           nonredundant cDNA library. Genome Res. . 11 (2), 281-289 (2001)
            Please visit our web site (http://genome.gsc.riken.go.jp) for
           further details.
            e mouse tissues.
                    Location/Qualifiers
FEATURES
                    1. .393
     source
                    /organism="Mus musculus"
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                    /db xref="taxon:10090"
                    /clone="G630022C22"
                    /sex="male"
                    /tissue type="intestinal mucosa"
                    /dev stage="adult"
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ORIGIN
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                                 Pred. No. 0.0045;
  Best Local Similarity
                         76.2%;
                                                                           2;
                                                      Indels
                                                                    Gaps
 Matches
           80; Conservative
                                0; Mismatches
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            4 GTAGGTGAGATCTCTGACCTCCAGAGTG---TTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
              318 GTATGTTAGATCTCTTACCTCCATAGTGGTTTGAGCTTACCAGCTCTAGGTTAAGTACAG 259
Db
           61 ACTGTTGTCACTTTCCGAGGAGAAC---AAGCTGTCCTGGAGGCC 102
Qy
              258 ACTGTTGTCACTTTCCTAGGAGTAAGCAAGGCTGTCCTGGAGGGC 214
Db
RESULT 11
BB605863/c
                                                               EST 05-DEC-2000
                                                      linear
                                    306 bp
                                              mRNA
LOCUS
            BB605863
           BB605863 RIKEN full-length enriched, 0 day neonate lung Mus
DEFINITION
            musculus cDNA clone E030013I04 5', mRNA sequence.
            BB605863
ACCESSION
VERSION
            BB605863.1 GI:11557265
KEYWORDS
            EST
            Mus musculus (house mouse)
SOURCE
  ORGANISM
           Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
               (bases 1 to 306)
REFERENCE
            Aizawa, K., Akahira, S., Akimura, T., Arai, A., Arakawa, T.,
  AUTHORS
            Carninci, P., Hanagaki, T., Hayatsu, N., Hiraoka, T., Hirozane, T.,
            Hodoyama, Y., Imotani, K., Ishii, Y., Itoh, M., Izawa, M., Kawai, J.,
            Kojima, Y., Konno, H., Kusakabe, M., Matsuyama, T., Miyazaki, A.,
            Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Okazaki, Y.,
            Okido, T., Owa, C., Sakai, C., Sakai, K., Sasaki, D., Sato, K.,
            Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T., Sogabe, Y.,
            Suzuki, H., Tagawa, A., Takahashi, F., Tanaka, T., Toya, T.,
            Watahiki, A., Yamamura, T., Yasunishi, A., Yoshida, K., Yoshiki, A.,
            Muramatsu, M. and Hayashizaki, Y.
            RIKEN Mouse ESTs (Aizawa, K. et al. 2000)
  TITLE
```

Computer-based methods for the mouse full-length cDNA

```
JOURNAL
           Unpublished (2000)
COMMENT
           Contact: Yoshihide Hayashizaki
           Laboratory for Genome Exploration Research Group, RIKEN Genomic
           Sciences Center(GSC), Yokohama Institute
           The Institute of Physical and Chemical Research (RIKEN)
           1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
           Tel: 81-45-503-9222
           Fax: 81-45-503-9216
           Email: genome-res@gsc.riken.go.jp,
           URL:http://genome.gsc.riken.go.jp/
           Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagaoka, S.,
           Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
           Thermostabilization and thermoactivation of thermolabile enzymes by
           trehalose and its application for the synthesis of full length
           cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
            Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.,
           Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M.,
           Okazaki, Y. and Hayashizaki, Y.
            Automated filtration-based high-throughput plasmid preparation
           system. Genome Res. 9 (5), 463-470 (1999)
            Carninci, P. and Hayashizaki, Y.
            High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
           19-44 (1999)
            Please visit our web site (http://genome.rtc.riken.go.jp) for
           further details.
                    Location/Qualifiers
FEATURES
    source
                    1. .306
                    /organism="Mus musculus"
                    /mol type="mRNA"
                    /db xref="taxon:10090"
                    /clone="E030013I04"
                    /tissue type="lung"
                    /dev stage="0 day neonate"
                    /lab host="DH10B"
                    /clone lib="RIKEN full-length enriched, 0 day neonate
                    /note="Site 1: SalI; Site_2: BamHI; cDNA library was
                    prepared and sequenced in Mouse Genome Encyclopedia
                    Project of Genome Exploration Research Group in Riken
                    Genomic Sciences Center and Genome Science Laboratory in
                    RIKEN. Division of Experimental Animal Research in Riken
                    contributed to prepare mouse tissues. 1st strand cDNA was
                    primed with a primer [5'
                    prepared by using trehalose thermo-activated reverse
                    transcriptase and subsequently enriched for full-length by
                    cap-trapper. Second strand cDNA was prepared with the
                    primer adapter of sequence [5'
                    was cleaved with BamHI and XhoI. Vector: a modified
                    pBluescript KS(+) after bulk excision from Lambda FLC I."
ORIGIN
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Query Match 39.0%; Score 39.8; DB 10; Length 306; Best Local Similarity 74.6%; Pred. No. 0.032; Matches 50; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

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Qу
                304 ACCCACCTCCGTAGTTGCAGTTCCAACTCTTGTCAATTTCCGAGGAGCACCACCTATCCA 245
Db
          96 GGAGGCC 102
QУ
             1111
         244 GGAGCCC 238
Db
RESULT 12
BM735433
LOCUS
           BM735433
                                    522 bp
                                              mRNA
                                                      linear
                                                               EST 01-MAR-2002
DEFINITION
           MONO1 20 F01.g1 A005 Monocytes (MONO1) Equus caballus cDNA, mRNA
            sequence.
ACCESSION
           BM735433
VERSION
           BM735433.1 GI:19056766
           EST.
KEYWORDS
SOURCE
            Equus caballus (horse)
  ORGANISM Equus caballus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Perissodactyla; Equidae; Equus.
REFERENCE
              (bases 1 to 522)
           Vandenplas, M.L., Cordonnier-Pratt, M.-M., Sudman, M.L., Wentzel, V.E.,
  AUTHORS
            Gingle, A.R., Pratt, L.H. and Moore, J.N.
           An EST database from equine (Equus caballus) monocytes
  TITLE
            Unpublished (2001)
  JOURNAL
            Contact: Cordonnier-Pratt MM
COMMENT
            Laboratory for Genomics and Bioinformatics
            The University of Georgia, Department of Plant Biology
            Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
            Tel: 706 542 1860
            Fax: 706 583 0210
            Email: mmpratt@uga.edu
            Sequences have been trimmed to exclude PolyA, vector and regions
            below Phred quality 16. The threshold for high quality sequence is
            20. Three-prime sequences, which are obtained with PolyTMix or T7
            sequencing primer, are presented as the reverse complement.
            Seq primer: T7
            High quality sequence start: 43
            High quality sequence stop: 522
            POLYA=Yes.
FEATURES
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                     1. .522
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                     /mol type="mRNA"
                     /db xref="taxon:9796"
                     /cell type="Isolated peripheral blood monocytes stimulated
                     with E. coli lipopolysaccharide"
                     /clone lib="Monocytes (MONO1)"
                     /note="Vector: pBluescript SK(-) from Lambda ZapII;
                     Site 1: XhoI; Site 2: EcoRI; The library was made from
                     poly-A RNA in the cloning vector lambda ZAPII. Clones to
                     be sequenced were prepared by mass excision."
ORIGIN
                         32.5%; Score 33.2; DB 12; Length 522;
  Query Match
  Best Local Similarity 67.1%; Pred. No. 5.7;
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0; Mismatches
                                                 23: Indels
                                                                0;
                                                                    Gaps
                                                                            0;
 Matches
           47; Conservative
            6 AGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGT 65
Qy
              11 1 1 11 1
                              200 AGTTCAAATATTGCCTGTCCAGAGAGGTTGTCCGACCACTGTAGCTGAAGCAGCGTCTCC 259
Db
          66 TGTCACTTTC 75
Qу
               260 AGTCACTTTC 269
Db
RESULT 13
AA524439/c
LOCUS
           AA524439
                                    597 bp
                                              mRNA
                                                      linear
                                                               EST 05-AUG-1997
           ng44f07.sl NCI CGAP Co3 Homo sapiens cDNA clone IMAGE:937669 3'
DEFINITION
            similar to qb:M28668 CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE
            REGULATOR (HUMAN);, mRNA sequence.
ACCESSION
           AA524439
VERSION
           AA524439.1 GI:2265367
KEYWORDS
            EST.
           Homo sapiens (human)
SOURCE
  ORGANISM
           Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              (bases 1 to 597)
REFERENCE
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
  AUTHORS
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
  TITLE
            Tumor Gene Index
            Unpublished (1997)
  JOURNAL
            Contact: Robert Strausberg, Ph.D.
COMMENT
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck,
            M.D., Ph.D.
             cDNA Library Preparation: M. Bento Soares, Ph.D.
             cDNA Library Arraying: Greg Lennon, Ph.D.
             DNA Sequencing by: Washington University Genome Sequencing Center
             Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Insert Length: 698
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            Seg primer: -40ml3 fwd. ET from Amersham
            High quality sequence stop: 322.
                     Location/Qualifiers
FEATURES
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                     /lab host="DH10B"
                     /clone lib="NCI CGAP Co3"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                     polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
                     was prepared from 12 pooled bulk tumor samples and primed
                     with a Not I - oligo(dT) primer. Double-stranded cDNA was
                     ligated to Eco RI adaptors (Pharmacia), digested with Not
```

I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. "

ORIGIN

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31.6%; Score 32.2; DB 9; Length 597;
 Ouery Match
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                         63.6%; Pred. No. 13;
                                                                           0;
                                                               0; Gaps
 Matches
           49; Conservative
                                0; Mismatches
                                                 28; Indels
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Qy
             112 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 53
Dh
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Qγ
             Dh
          52 TTCTCAGTTTTCCTGGA 36
RESULT 14
BX506811
                                                               EST 04-SEP-2003
                                    752 bp
                                             mRNA
                                                      linear
LOCUS
           BX506811
           DKFZp779M191 rl 779 (synonym: hnccl) Homo sapiens cDNA clone
           DKFZp779M191 5', mRNA sequence.
ACCESSION
           BX506811
           BX506811.1 GI:32047420
VERSION
KEYWORDS
           EST.
SOURCE
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           Homo sapiens
  ORGANISM
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              (bases 1 to 752)
REFERENCE
           Poustka, A., Albert, R., Moosmayer, P., Schupp, I., Wellenreuther, R.,
  AUTHORS
           Mewes, H.W., Weil, B., Amid, C., Osanger, A., Fobo, G., Han, M. and
           Wiemann, S.
           EST (Poustka, A., Albert, R., Moosmayer, P., Schupp, I.,
  TITLE
           Wellenreuther, R., et al.)
           Unpublished (2003)
  JOURNAL
           Contact: MIPS
COMMENT
           MIPS
           Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
           This is the 5' sequence of the clone insert
           Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
            Research Center (DKFZ); Email s.wiemann@dkfz- heidelberg.de;
            sequenced by DKFZ (German Cancer Research Center,
            Heidelberg/Germany) within the cDNA sequencing consortium of the
            German Genome Project.
           No s1 sequence available.
            This clone (DKFZp779M191) is available at the RZPD in Berlin.
            Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
            Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
                     Location/Qualifiers
FEATURES
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                     /db xref="taxon:9606"
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                     /tissue type="liver"
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/dev stage="fetal"
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  Query Match
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                          63.6%;
                                  Pred. No. 15;
                                                                  0;
                                                                      Gaps
                                                                              0;
            49; Conservative
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                                                  28;
                                                       Indels
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Qу
                              11 1 1111 11
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Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              363 TTCTCAGTTTTCCTGGA 379
Db
RESULT 15
AY399795
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                                                                 GSS 15-DEC-2003
DEFINITION
            Homo sapiens CFTR gene, VIRTUAL TRANSCRIPT, partial sequence,
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ACCESSION
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VERSION
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KEYWORDS
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SOURCE
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REFERENCE
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            Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
  AUTHORS
            Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
            Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
            Adams, M.D. and Cargill, M.
            Inferring nonneutral evolution from human-chimp-mouse orthologous
  TITLE
            gene trios
  JOURNAL
            Science 302 (5652), 1960-1963 (2003)
   PUBMED
            14671302
REFERENCE
            2 (bases 1 to 4443)
            Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
  AUTHORS
            Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
            Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
            Adams, M.D. and Cargill, M.
  TITLE
            Direct Submission
            Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
  JOURNAL
            Rockville, MD 20850, USA
            This sequence was made by sequencing genomic exons and ordering
COMMENT
            them based on alignment.
FEATURES
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                     /mol type="genomic DNA"
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ORIGIN

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                                                                            0;
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
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QУ
              11 | 1111 | 11
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Db
          65 TTGTCACTTTCCGAGGA 81
Qу
              1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 16
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                                    500 bp
                                              mRNA
                                                      linear
                                                               EST 06-DEC-2000
LOCUS
           WHE2306 H10 O20ZS Wheat pre-anthesis spike cDNA library Triticum
DEFINITION
           aestivum cDNA clone WHE2306 H10 O20, mRNA sequence.
ACCESSION
           BF483989
VERSION
           BF483989.1 GI:11567278
KEYWORDS
           EST.
           Triticum aestivum (bread wheat)
SOURCE
           Triticum aestivum
  ORGANISM
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Pooideae; Triticeae; Triticum.
              (bases 1 to 500)
REFERENCE
  AUTHORS
           Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D.,
           Han, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J.,
            Seaton, C.L. and Tong, J.C.
  TITLE
            The structure and function of the expressed portion of the wheat
            genomes - Pre-anthesis spike cDNA library
  JOURNAL
            Unpublished (2000)
            Contact: Olin Anderson
COMMENT
            US Department of Agriculture, Agriculture Research Service, Pacific
            West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
            Tel: 5105595773
            Fax: 5105595818
            Email: oandersn@pw.usda.gov
            Sequence have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seq primer: Stratagene SK primer.
FEATURES
                    Location/Qualifiers
                     1. .500
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                     /mol type="mRNA"
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                     /dev stage="Adult plant"
                     /lab host="E. coli SOLR"
                     /clone lib="Wheat pre-anthesis spike cDNA library"
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/note="Vector: Lambda Uni-ZAP XR, excised phagemid; Site_1: EcoRI; Site_2: XhoI; Plants were grown in the greenhouse. Whole spike with awns trimmed, white, green and yellow anther were collected and total RNA, and poly(A) RNA were prepared, a cDNA library was made, and the cDNA clones were in vivo excised to give pBluescript phagemids in the TJ Close lab (Choi, Close, Fenton) at the University of California, Riverside. Plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors)."

ORIGIN

Query Match 30.6%; Score 31.2; DB 10; Length 500; Best Local Similarity 57.0%; Pred. No. 25; 57; Conservative 0; Mismatches 43; Indels 0; Gaps 0; 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qу 1 111 | | | | | Db 389 CTGGTGCCCGCAATCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 330 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100 Qу Db 329 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGATG 290

RESULT 17 BE404165/c

LOCUS BE404165 693 bp mRNA linear EST 21-JUL-2000

DEFINITION WHE1201 H12 O23ZS Wheat etiolated seedling root cDNA library

Triticum aestivum cDNA clone WHE1201_H12_O23, mRNA sequence.

ACCESSION BE404165

VERSION BE404165.1 GI:9363633

KEYWORDS EST.

SOURCE Triticum aestivum (bread wheat)

ORGANISM Triticum aestivum

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

Pooideae; Triticeae; Triticum.

REFERENCE 1 (bases 1 to 693)

AUTHORS Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D.,

Han, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J.,

Seaton, C.L. and Tong, J.C.

TITLE The structure and function of the expressed portion of the wheat

genomes

JOURNAL Unpublished (2000)

COMMENT Contact: Olin Anderson

US Department of Agriculture, Agriculture Research Service, Pacific

West Area, Western Regional Research Center 800 Buchanan Street, Albany, CA 94710, USA

Tel: 5105595773 Fax: 5105595818

Email: oandersn@pw.usda.gov

Sequence have been trimmed to remove vector sequence and low

quality sequence with phred score less than 20

Seq primer: Strategene SK primer.

FEATURES Location/Qualifiers

source 1. .693

/organism="Triticum aestivum" /mol type="mRNA" /cultivar="Chinese Spring" /db xref="taxon:4565" /clone="WHE1201 H12 023" /tissue type="Root" /dev stage="Five day old etiolated seedling" /lab host="E. coli SOLR" /clone lib="Wheat etiolated seedling root cDNA library" /note="Vector: Lambda Uni-ZAP XR, excised phagemid; Site 1: EcoRI; Site_2: XhoI; Seeds were surface-sterilized, germinated and grown aseptically in the dark at room temperature on filter paper with water, nystatin and cefotaxime in covered crystallization dishes. Roots were harvested. The tissue, total RNA, and poly(A) RNA were prepared, a cDNA library was made, and the cDNA clones were in vivo excised to give pBluescript phagemids in the TJ Close lab (Choi, Close, Fenton) at the University of California, Riverside. Plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors)."

ORIGIN

Query Match 30.6%; Score 31.2; DB 10; Length 693; Best Local Similarity 57.0%; Pred. No. 29; 57; Conservative 0; Mismatches 43; Indels Gaps 0; 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qу 1 111 Db 319 CTGGTGCCCGCAATCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 260 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100 Qу 259 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGATG 220 Db

RESULT 18 BH295020/c

LOCUS BH295020 648 bp DNA linear GSS 30-NOV-2001 DEFINITION CH230-44L24.TJ CHORI-230 Segment 1 Rattus norvegicus genomic clone

CH230-44L24, genomic survey sequence.

ACCESSION BH295020

VERSION BH295020.1 GI:17207428

KEYWORDS GSS.

SOURCE Rattus norvegicus (Norway rat)

ORGANISM Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE 1 (bases 1 to 648)

AUTHORS Zhao, S., Shetty, J., Shatsman, S., Tsegaye, G., Geer, K.,

Shvartsbeyn, A., Gebregeorgis, E., Overton, L., Russell, D., Chen, D.,

Riggs, F., de Jong, P. and Fraser, C.M.

TITLE Rat BAC End Sequences from Library CHORI-230 EcoRI segment

JOURNAL Unpublished (1999)

COMMENT Other_GSSs: CH230-44L24.TV

Contact: Shaying Zhao

```
Department of Eukaryotic Genomics
           The Institute for Genomic Research
           9712 Medical Center Dr., Rockville, MD 20850, USA
           Tel: 301 838 0200
           Fax: 301 838 0208
           Email: szhao@tigr.org
           Clones are derived from the rat BAC library CHORI-230
           (http://www.chori.org/bacpac/rat230.htm). For BAC library
           availability, please contact Pieter de Jong (pdejong@mail.cho.org).
           Clones may be purchased from BACPAC Resources
           (http://www.chori.org/bacpac/or ering information.htm). BAC end
           page: http://www.tigr.org/tdb/bac ends/rat/bac end intro.html
           Plate: 44 row: L column: 24
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                    /clone="CH230-44L24"
                    /sex="Female"
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                    /clone lib="CHORI-230 Segment 1"
                    /note="Vector: pTARBAC2.1; Site 1: EcoRI; Site 2: EcoRI;
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                                0: Mismatches
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                                                     Indels
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QУ
             355 GTGAGTTCTGTCACCTGCAAGGCAAAGGCCAAAGCAGAAAAACTAAACTACAGAGGGAAG 296
Db
          68 TCACTTTCCGAGGAGAACAAGCTGTCC 94
Qу
                         Dh
         295 TCTCTTTCTTGTGAGAACATTCTCACC 269
RESULT 19
AV277244/c
                                                              EST 05-NOV-1999
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                                                     linear
LOCUS
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           AV277244 RIKEN full-length enriched, adult male testis (DH10B) Mus
DEFINITION
           musculus cDNA clone 4932441F23 3', mRNA sequence.
           AV277244
ACCESSION
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VERSION
KEYWORDS
           EST.
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SOURCE
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            1 (bases 1 to 238)
REFERENCE
           Konno, H., Aizawa, K., Akahira, S., Akiyama, J., Carninci, P., Endo, T.,
 AUTHORS
```

Fukuda, S., Fukunishi, Y., Hara, A., Hayatsu, N., Hirozane, T., Hori, F., Ishii, Y., Ishikawa, T., Itoh, M., Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N., Kojima, Y., Koya, S., Kusakabe, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Owa, C., Ozawa, Y., Saito, H., Sano, M., Sato, K., Shibata, K., Shibata, Y., Shiqemoto, Y., Shiraki, T., Soqabe, Y., Sugahara, Y., Suzuki, H., Suzuki, H., Takahashi, F., Tateno, M., Tominaga, N., Tsunoda, Y., Watahiki, A., Watanabe, S., Yamamura, T., Yasunishi, A., Yokota, T., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y. RIKEN Mouse ESTs (Konno, H., et al. 1999) Unpublished (1999) Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/ Sasaki, N., Izawa, M., Watahiki, M., Ozawa, K., Tanaka, T., Yoneda, Y., Matsuura, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki,Y. Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y. Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999) Carninci, P. and Hayashizaki, Y. High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999) Please visit our web site (http://genome.rtc.riken.go.jp) for further details. Location/Qualifiers 1. .238 source /organism="Mus musculus" /mol type="mRNA" /strain="C57BL/6J" /db xref="taxon:10090" /clone="4932441F23" /sex="male" /tissue type="testis" /dev stage="adult" /lab host="DH10B" /clone lib="RIKEN full-length enriched, adult male testis (DH10B)" /note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5'

> prepared by using trehalose thermo-activated reverse

TITLE JOURNAL

COMMENT

FEATURES

ORIGIN

Score 30.4; DB 9; Length 238; Query Match 29.8%; Best Local Similarity 59.1%; Pred. No. 30; 52; Conservative Matches 0; Mismatches 36; Indels 0: Gaps 0; 14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73 Qу | | | | | Π Db 222 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATAAAGTTGCTCTTGAAAACACTT 163 74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101 Qу 1 111 Db 162 TTCGATAAGAACAATCTGTTCTTGTAGC 135 RESULT 20 AA706660 LOCUS AA706660 294 bp mRNA linear EST 24-DEC-1997 ag90h11.rl Stratagene hNT neuron (#937233) Homo sapiens cDNA clone DEFINITION IMAGE: 1141797 5', mRNA sequence. ACCESSION AA706660 AA706660.1 GI:2716578 VERSION KEYWORDS EST. SOURCE Homo sapiens (human) Homo sapiens ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 294) REFERENCE Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S., AUTHORS Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R. TITLE WashU-NCI human EST Project JOURNAL Unpublished (1997) Contact: Wilson RK COMMENT Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Seq primer: -28m13 rev1 ET from Amersham High quality sequence stop: 281. **FEATURES** Location/Qualifiers 1. .294 source /organism="Homo sapiens" /mol type="mRNA" /db xref="taxon:9606"

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                    GAATTCGGCACGAG 3' ~3' adaptor sequence: 5'
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ORIGIN
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 Query Match
 Best Local Similarity
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                                 Pred. No. 34;
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                                0; Mismatches
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                                                     Indels
                                                                  Gaps
                                                                          0;
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Qу
             1 11
                                                          Db
          64 GTTGTCACTTTCCGAGGAGAACAAGCTG 91
QУ
                    \Box\Box\Box
                           Db
         112 TTGGGGGCTTTGAAAGGAGAACAGCGTG 139
RESULT 21
BG980021/c
LOCUS
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                                    295 bp
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DEFINITION
ACCESSION
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VERSION
           BG980021.1 GI:14382756
KEYWORDS
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SOURCE
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 ORGANISM
           Homo sapiens
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REFERENCE
              (bases 1 to 295)
           Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,
  AUTHORS
           Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,
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           Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V.,
           O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and
           Simpson, A.J.
           Shotgun sequencing of the human transcriptome with ORF expressed
 TITLE
           sequence tags
           Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
  JOURNAL
           20202663
  MEDLINE
           10737800
   PUBMED
           Contact: Simpson A.J.G.
COMMENT
           Laboratory of Cancer Genetics
           Ludwig Institute for Cancer Research
           Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
           Brazil
           Tel: +55-11-2704922
           Fax: +55-11-2707001
           Email: asimpson@ludwig.org.br
           This sequence was derived from the FAPESP/LICR Human Cancer Genome
           Project. This entry can be seen in the following URL
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/lab host="SOLR (kanamycin resistant)"

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                     derived from ORESTES PCR (U.S. Letters Patent application
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                     profiles into the pUC 18 vector. Reverse transcription of
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ORIGIN
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Qy
                         294 GAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTGTTCTC 235
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Qу
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Db
RESULT 22
AV254401/c
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                                                       linear
                                                                EST 24-OCT-2001
                                     986 bp
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LOCUS
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DEFINITION
            musculus cDNA clone 4921509J16 3', mRNA sequence.
ACCESSION
            AV254401
            AV254401.2 GI:16388054
VERSION
KEYWORDS
            EST.
            Mus musculus (house mouse)
SOURCE
           Mus musculus
  ORGANISM
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            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
              (bases 1 to 986)
REFERENCE
  AUTHORS
            Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T.,
            Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J.,
            Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K.,
            Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,
            Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
            Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
            Takeda, Y., Tanaka, T., Toya, T., Muramatsu, M. and Hayashizaki, Y.
            RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
  TITLE
            Unpublished (2001)
  JOURNAL
            On Nov 4, 1999 this sequence version replaced gi:6241860.
COMMENT
```

Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/ Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. . 10 (10), 1617-1630 (2000) wagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. . 10 (11), 1757-1771 (2000) Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y. Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. . 11 (2), 281-289 (2001) Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I., Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Havashizaki,Y. Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001) Please visit our web site (http://genome.gsc.riken.go.jp/) for further details. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. Location/Qualifiers 1. .986 /organism="Mus musculus" /mol type="mRNA" /strain="C57BL/6J" /db xref="taxon:10090" /clone="4921509J16" /sex="male" /tissue type="testis" /dev stage="adult" /lab host="DH10B" /clone lib="RIKEN full-length enriched, adult male testis (DH10B)"

/note="Site_1: SalI; Site_2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was

FEATURES

source

prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5' was cloned into the XhoI and BamHI sites. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI." ORIGIN Query Match 29.8%; Score 30.4; DB 9; Length 986; Best Local Similarity 59.1%; Pred. No. 64; 52: Conservative 0; Mismatches 36; Indels Gaps 0; 14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73 Qу 1 1111 11 Db 375 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 316 74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101 Qy -Db 315 TCCGATGAGAGCGATCTGTTCTTGTAGC 288 RESULT 23 CC921947/c 746 bp DNA linear GSS 08-AUG-2003 LOCUS CC921947 DEFINITION t060j23ba.fl TAMBT Bos taurus genomic clone t060j23ba, genomic survey sequence. ACCESSION CC921947 CC921947.1 GI:33555987 VERSION KEYWORDS GSS. Bos taurus (cow) SOURCE ORGANISM Bos taurus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovoidea; Bovidae; Bovinae; Bos. REFERENCE 1 (bases 1 to 746) AUTHORS Lin, S., Najar, F.Z., Adelson, D., Gill, C.A. and Roe, B.A. Bovine BAC End Sequences from Library TAMBT TITLE Unpublished (2003) JOURNAL Contact: Bruce A. Roe COMMENT Advanced Center for Genome Technology University of Oklahoma Department of Chemistry and Biochemistry 620 Parrington Oval, Room 208, Norman, OK 73019, USA Tel: 405 325 4912 Fax: 405 325 7762 Email: broe@ou.edu Class: BAC ends High quality sequence start: 40 High quality sequence stop: 739. Location/Qualifiers **FEATURES** 1. .746 source /organism="Bos taurus" /mol type="genomic DNA" /strain="Angus bull T A M U Shoshone Y6 11519666"

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ORIGIN
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Qу
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             Db
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DEFINITION
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VERSION
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SOURCE
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              (bases 1 to 551)
REFERENCE
           Sanaka, E., Hori, H., Naruse, K., Mitani, H. and Tanaka, M.
 AUTHORS
 TITLE
           Medaka EST analysis
 JOURNAL
           Unpublished (2001)
           Contact: Emi Sanaka
COMMENT
           Department of Biological Sciences
           Graduate School of Science, Nagoya University
           Furo-cho, Chikusa-ku, Nagoya 464-8602, Japan
           Tel: 81-52-789-2973
           Fax: 81-52-789-2974
           Email: sanaka@bio.nagoya-u.ac.jp
           This clone was isolated from Medaka eye cDNA library (SNK01) 5'end.
FEATURES
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Qу
                        372 CTGGCAGGGTTGATTTCTAAGATAAAAAGTGTAGGACTGATTACTGTTGAAGAAGAAGAAA 431
Db
          61 ACTGTTGTCACTT 73
Qу
              | | |
          432 AGTCCAGGTGCTT 444
Db
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DEFINITION
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           BF473385
ACCESSION
VERSION
           BF473385.1 GI:11542567
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SOURCE
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           Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
           Pooideae; Triticeae; Triticum.
               (bases 1 to 444)
REFERENCE
           1
           Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D.,
  AUTHORS
           Han, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J.,
            Seaton, C.L. and Tong, J.C.
           The structure and function of the expressed portion of the wheat
  TITLE
            genomes - 5-15 DAP spike cDNA library
  JOURNAL
           Unpublished (2000)
COMMENT
           Contact: Olin Anderson
           US Department of Agriculture, Agriculture Research Service, Pacific
           West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
           Tel: 5105595773
            Fax: 5105595818
            Email: oandersn@pw.usda.gov
            Sequence have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seq primer: Stratagene SK primer.
                     Location/Qualifiers
FEATURES
                     1. .444
     source
                     /organism="Triticum aestivum"
                     /mol type="mRNA"
                     /cultivar="Chinese Spring"
                     /db xref="taxon:4565"
                     /clone="WHE0923 H02 P03"
                     /tissue type="Spike"
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/dev stage="adult"

/dev stage="Adult plant" /lab host="E. coli SOLR" /clone lib="Wheat 5-15 DAP spike cDNA library" /note="Vector: Lambda Uni-ZAP XR, excised phagemid; Site 1: EcoRI; Site 2: XhoI; Plants were grown in the greenhouse. Spikes at 5, 10 and 15 DAP were harvested, total RNA and poly(A) RNA were prepared, a cDNA library was made, and the cDNA clones were in vivo excised to give pBluescript phagemids in the TJ Close lab (Choi, Close, Fenton) at the University of California, Riverside. Plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors)." 29.0%; Score 29.6; DB 10; Length 444; 56.0%; Pred. No. 76; Best Local Similarity 56; Conservative 0; Mismatches 44; Indels 0; Gaps 0; 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 1 323 CTGGTGCCCGCAATCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 264 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100 263 CTCCGTGTCGCTCACCTCCCAGAACATGTCGTCGTCGATG 224 EST 30-MAY-2002 B0467131 483 bp mRNA linear HS02L11r HS Hordeum vulgare subsp. vulgare cDNA clone HS02L11 5-PRIME, mRNA sequence. B0467131 BQ467131.1 GI:21274913 Hordeum vulgare subsp. vulgare Hordeum vulgare subsp. vulgare Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Hordeum. (bases 1 to 483) Zhang, H., Potokina, E., Michalek, W., Weschke, W., Stein, N. and Graner, A. Barley ESTs from germinating seeds Unpublished (2002) Contact: Stein Nils Molecular Markers Group, Department Genbank Institute of Plant Genetics and Crop Plant Research (IPK) Corrensstr. 3, 06466, Gatersleben, Germany Tel: 039482-5522 Fax: 039482-5595 Email: stein@ipk-gatersleben.de Insert Length: 483 Std Error: 0.00 Plate: 2 row: L column: 11 Seq primer: M13rev.

ORIGIN

QУ

Db

Qу

Db

RESULT 26 B0467131/c

DEFINITION

ORGANISM

EST.

Location/Qualifiers

ACCESSION

REFERENCE

TITLE

FEATURES

AUTHORS

JOURNAL COMMENT

VERSION KEYWORDS

SOURCE

LOCUS

Query Match

1. .483 source /organism="Hordeum vulgare subsp. vulgare" /mol type="mRNA" /cultivar="barke" /sub species="vulgare" /db xref="taxon:112509" /clone="HS02L11" /tissue type="embryo + scutellum" /dev stage="0-16 hours after imbibition" /lab host="XL10-Gold" /clone lib="HS" /note="Vector: pBluescript SK+; Site 1: EcoRI (5'-end of cDNA); Site 2: XhoI (3'-end of cDNA); Due to a cloning artefact caused by the kit, in most cases the EcoRI site is NOT present, as well as the EcoRIadapter used for cloning. To excise the insert, restriction sites upstream EcoRI should be used (e.g. BamHI, SalI, PstI). NOTE: Also due to the cloning system used Blue/white selection for recombinats is not 100% reliable." ORIGIN 29.0%; Score 29.6; DB 13; Length 483; Query Match Best Local Similarity 56.0%; Pred. No. 80; 56; Conservative 0; Mismatches 44; Indels 0; Gaps Matches 11 IIIIII61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100

1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qy 375 CTGGTGCCCGCAGTCCTCCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 316 Db Qv

315 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGACG 276

0;

RESULT 27 CD912921/c

Db

EST 14-JUL-2003 CD912921 mRNA linear LOCUS 518 bp G550.116E20F010525 G550 Triticum aestivum cDNA clone G550116E20, DEFINITION mRNA sequence.

ACCESSION CD912921

VERSION CD912921.1 GI:32687245

KEYWORDS EST.

SOURCE Triticum aestivum (bread wheat)

ORGANISM Triticum aestivum

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

Pooideae; Triticeae; Triticum.

(bases 1 to 518) REFERENCE

Genoplante. AUTHORS

Genoplante, a major partnership french program in plant genomics TITLE

JOURNAL Unpublished (2003) COMMENT Contact: Genoplante

Genoplante

93, rue Henri Rochefort 91025 EVRY CEDEX France

Tel: 33 1 69 47 54 00 Fax: 33 1 69 47 54 10

This sequence has been generated in the framework of the french

```
plant genomics programme 'Genoplante' (http://www.genoplante.com
           and http://genoplante-info.infobiogen.fr).
FEATURES
                    Location/Qualifiers
    source
                    1. .518
                    /organism="Triticum aestivum"
                    /mol type="mRNA"
                    /cultivar="recital"
                    /db xref="taxon:4565"
                    /clone="G550116E20"
                    /tissue type="grain (550 degrees per day after
                    pollination)"
                    /clone lib="G550"
ORIGIN
 Query Match
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 Best Local Similarity
                         56.0%; Pred. No. 82;
           56; Conservative
                                0; Mismatches
                                                 44;
                                                      Indels
                                                                0;
                                                                    Gaps
                                                                            0;
 Matches
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
                          1 111 1
                                                          1 | | | | | | | | | | |
             \perp
          393 CTGGTGCCCGCAGTCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 334
Db
          61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100
Qy
                   333 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGATG 294
Db
RESULT 28
BG606129/c
                                    564 bp
                                                      linear
                                                               EST 17-APR-2001
LOCUS
           BG606129
                                              mRNA
           WHE2960 H03 006ZS Wheat dormant embryo cDNA library Triticum
           aestivum cDNA clone WHE2960 H03 006, mRNA sequence.
ACCESSION
           BG606129
           BG606129.1 GI:13656112
VERSION
KEYWORDS
           EST.
           Triticum aestivum (bread wheat)
SOURCE
  ORGANISM Triticum aestivum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Pooideae; Triticeae; Triticum.
REFERENCE
              (bases 1 to 564)
           Anderson, O.D., Chao, S., Chin, A., Close, T.J., Doherty, L.,
  AUTHORS
            Fenton, R.D., Lazo, G.R., Rausch, C.J., Walker-Simmons, M.K. and
            Wilson, C.
            The structure and function of the expressed portion of the wheat
  TITLE
            genomes - Dormant embryo cDNA library
           Unpublished (2001)
  JOURNAL
COMMENT
            Contact: Olin Anderson
            US Department of Agriculture, Agriculture Research Service, Pacific
            West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
            Tel: 5105595773
            Fax: 5105595818
            Email: oandersn@pw.usda.gov
            Sequence have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seq primer: Stratagene SK primer.
```

FEATURES source

Location/Qualifiers

1. .564

/organism="Triticum aestivum"

/mol type="mRNA" /cultivar="Brevor" /db xref="taxon:4565" /clone="WHE2960 H03 006" /tissue type="Seed embryo"

/dev stage="Mature seed" /lab_host="E. coli SOLR"

/clone lib="Wheat dormant embryo cDNA library" /note="Vector: Lambda Uni-ZAP XR, excised phagemid; Site 1: EcoRI; Site 2: XhoI; Plants were grown to seed maturity under conditions favoring seed dormancy (L. Dohery at K. Walker Simmons lab, Washington State University, Pullman, WA). Embryos were cut from mature dormant seed (Doherty). Total RNA was prepared from these embryos, polyA was purified, a cDNA library was made, and the cDNA clones were in vivo excised to give pBluescript phagemids in the TJ Close lab at the University of California, Riverside (Chin, Fenton). Plasmid DNA preparations and DNA sequencing were performed in the OD

Anderson lab (all other authors)."

ORIGIN

29.0%; Score 29.6; DB 12; Length 564; Query Match 56.0%; Pred. No. 86; Best Local Similarity 0; 56; Conservative Gaps 0; Mismatches 44; Indels

1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qv 1 111 1 11 11111 1 111 336 CTGGTGCCCGCAGTCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 277 Db

61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100 Qy 1111 11 11 11 11111 1 111 1 11 1

276 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGATG 237 Db

RESULT 29 BM377546/c

EST 23-JUL-2002 BM377546 661 bp mRNA linear LOCUS EBem04 SQ003 H10 R embryo, 12 DPA, no treatment, cv Optic, EBem04 DEFINITION Hordeum vulgare subsp. vulgare cDNA clone EBem04 SQ003 H10 5', mRNA sequence.

BM377546 ACCESSION

BM377546.2 GI:21933449 VERSION

KEYWORDS EST.

Hordeum vulgare subsp. vulgare SOURCE Hordeum vulgare subsp. vulgare ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

Pooideae; Triticeae; Hordeum.

1 (bases 1 to 661) REFERENCE

Hedley, P., Liu, H., Caldwell, D., McCallum, N., Mudie, S., Cardle, L., AUTHORS

Ramsay, L., Machray, G., Marshall, D.F.M. and Waugh, R.

Development of Barley Transcriptome Resources TITLE

Unpublished (2001) JOURNAL

```
On Jan 10, 2002 this sequence version replaced gi:18120936.
COMMENT
           Contact: Waugh R, Marshall DF
           Genome Dynamics/Computational Biology
            Scottish Crop Research Institute
            Invergowrie, Dundee, DD2 5DA, Scotland, UK
           Tel: 00 44 1382 562731
            Fax: 00 44 1382 562426
            Email: est@scri.sari.ac.uk
           All sequence has a Phred quality score of 20 or over
            Seq primer: M13 reverse.
FEATURES
                     Location/Qualifiers
                     1. .661
     source
                     /organism="Hordeum vulgare subsp. vulgare"
                     /mol type="mRNA"
                     /cultivar="Optic"
                     /sub species="vulgare"
                     /db xref="taxon:112509"
                     /clone="EBem04_SQ003 H10"
                     /tissue type="embryo"
                     /dev_stage="12 DPA"
                     /lab host="DH10B"
                     /clone lib="embryo, 12 DPA, no treatment, cv Optic,
                     /note="Vector: pSPORT1; Site 1: Sal I; Site 2: Not I;
                     Non-normalised library, directionally cloned into pSPORT1.
                     Derived from embryos dissected from developing grains (12
                     days post anthesis) in glasshouse grown barley plants.
                     Developed as part of the barley transcriptome resources of
                     BBSRC/SEERAD funded cereal IGF (Investigating Gene
                     Function) project."
ORIGIN
                          29.0%; Score 29.6; DB 12;
                                                      Length 661;
  Query Match
                         56.0%; Pred. No. 94;
  Best Local Similarity
                                                                     Gaps
                                                                             0;
                                 0; Mismatches
                                                  44;
                                                      Indels
            56: Conservative
            1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qy
                                               11111 1 111
                       1
                          453 CTGGTGCCCGCAGTCCTCCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 394
Db
           61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100
Qу
                                 1111111 1 111 1 11 1
                   393 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGATG 354
Db
RESULT 30
BQ466828/c
                                                       linear
                                                                EST 30-MAY-2002
                                     697 bp
                                               mRNA
            BQ466828
LOCUS
            HS01L11T HS Hordeum vulgare subsp. vulgare cDNA clone HS01L11
DEFINITION
            5-PRIME, mRNA sequence.
ACCESSION
            BQ466828
            BQ466828.1 GI:21274610
VERSION
            EST.
KEYWORDS
            Hordeum vulgare subsp. vulgare
SOURCE
  ORGANISM Hordeum vulgare subsp. vulgare
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
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Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

```
Pooideae; Triticeae; Hordeum.
           1 (bases 1 to 697)
REFERENCE
           Zhang, H., Potokina, E., Michalek, W., Weschke, W., Stein, N. and
  AUTHORS
           Graner, A.
  TITLE
           Barley ESTs from germinating seeds
           Unpublished (2002)
  JOURNAL
           Contact: Stein Nils
COMMENT
           Molecular Markers Group, Department Genbank
           Institute of Plant Genetics and Crop Plant Research (IPK)
           Corrensstr. 3, 06466, Gatersleben, Germany
           Tel: 039482-5522
           Fax: 039482-5595
           Email: stein@ipk-gatersleben.de
           Insert Length: 697
                               Std Error: 0.00
           Plate: 1 row: L column: 11
           Seq primer: T3.
                    Location/Qualifiers
FEATURES
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                    /sub_species="vulgare"
                    /db xref="taxon:112509"
                    /clone="HS01L11"
                    /tissue type="embryo + scutellum"
                    /dev stage="0-16 hours after imbibition"
                    /lab host="XL10-Gold"
                    /clone lib="HS"
                    /note="Vector: pBluescript SK+; Site 1: EcoRI (5'-end of
                    cDNA); Site 2: XhoI (3'-end of cDNA); Due to a cloning
                    artefact caused by the kit, in most cases the EcoRI site
                    is NOT present, as well as the EcoRIadapter used for
                    cloning. To excise the insert, restriction sites upstream
                    EcoRI should be used (e.g. BamHI, SalI, PstI). NOTE: Also
                    due to the cloning system used Blue/white selection for
                    recombinats is not 100% reliable."
ORIGIN
                         29.0%; Score 29.6; DB 13; Length 697;
  Query Match
  Best Local Similarity
                         56.0%; Pred. No. 96;
                                0; Mismatches
                                                 44;
                                                      Indels
                                                                    Gaps
                                                                            0;
  Matches
           56; Conservative
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qy
                                1 111 1
                                                         1
                          \Box
          374 CTGGTGCCGCAGTCCTCCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAC 315
Db
          61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100
Qy
                                314 CTCCGTGTCGCTCACCTCCGAGAACATGTCGTCGTCGACG 275
Db
RESULT 31
BQ838111/c
                                                               EST 08-AUG-2002
                                    730 bp
                                                      linear
                                              mRNA
LOCUS
           BQ838111
           WHE2906 F08 K16ZS Wheat aluminum-stressed root tip cDNA library
DEFINITION
           Triticum aestivum cDNA clone WHE2906 F08 K16, mRNA sequence.
           BQ838111
ACCESSION
```

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BQ838111.1 GI:22142429
KEYWORDS
            EST.
SOURCE
            Triticum aestivum (bread wheat)
  ORGANISM
           Triticum aestivum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Pooideae; Triticeae; Triticum.
              (bases 1 to 730)
REFERENCE
            Anderson, O.D., Chao, S., Chin, A., Close, T.J., Gustafson, J.P.,
 AUTHORS
            Lazo, G.R., Rausch, C.J., Ross, K., Seaton, C.L. and Wilson, C.
  TITLE
            The structure and function of the expressed portion of the wheat
            genomes - Aluminum-stressed root tip cDNA library
            Unpublished (2001)
  JOURNAL
            Contact: Olin Anderson
COMMENT
            US Department of Agriculture, Agriculture Research Service, Pacific
            West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
            Tel: 5105595773
            Fax: 5105595818
            Email: oandersn@pw.usda.gov
            Sequences have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seq primer: SK primer.
                    Location/Qualifiers
FEATURES
                     1. .730
     source
                     /organism="Triticum aestivum"
                     /mol type="mRNA"
                     /cultivar="BH1146"
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                     /tissue type="Root tip at 1.0 to 1.5 mm stage"
                     /dev stage="Seedling"
                     /lab_host="E. coli SOLR"
                     /clone lib="Wheat aluminum-stressed root tip cDNA library"
                     /note="Vector: Lambda Uni-ZAP XR, excised phagemid;
                     Site 1: EcoRI; Site 2: XhoI; Plants were grown under
                     hydroponic conditions, root tips were excised and snap
                     frozen, total RNA was prepared at University of
                     Missouri (Ross, Gustafson). Poly(A) RNA was purified, a
                     cDNA library was made, and the cDNA clones were in vivo
                     excised to give pBluescript SK- phagemids in the TJ Close
                     lab (Chin and Close) at the University of California,
                     Riverside. Plasmid DNA preparations and DNA sequencing
                     were performed in the OD Anderson lab (all other
                     authors)."
ORIGIN
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  Query Match
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            56; Conservative
                                 0; Mismatches
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                                                       Indels
                                                                     Gaps
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Qy
                                 1 111 1
                                                          \perp
          146 CTGGTGCCCGCAGTCCTGCACCTCGATGGTGCACGCCTGGTCGAAGCAGATGCAGCACAG 87
Db
           61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100
QУ
                   1111 11 11 11 11 1 11 1 1 1
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VERSTON

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RESULT 32
CB360743
            CB360743
                                     429 bp
                                               mRNA
                                                       linear
                                                                 EST 10-NOV-2003
LOCUS
            ZF001-P00031-DPE-F-D B08 GISZF001 Danio rerio cDNA clone
DEFINITION
            IMAGE: 6903233 5' similar to fc20e06.yl Zebrafish WashU MPIMG EST
            Danio rerio cDNA clone IMAGE: 3721954 5', mRNA sequence.
ACCESSION
            CB360743
VERSION
            CB360743.1 GI:29005688
KEYWORDS
            EST.
            Danio rerio (zebrafish)
SOURCE
            Danio rerio
  ORGANISM
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
            Cypriniformes; Cyprinidae; Danio.
REFERENCE
            1 (bases 1 to 429)
  AUTHORS
            Mathavan, S., Wei, C., Thoreau, H., Chia, J.M. and Ruan, Y.
            Genome Institute of Singapore, Zebrafish EST Collection
  TITLE
            Unpublished (2003)
  JOURNAL
COMMENT
            Contact: Ruan Y
            Laboratory of Molecular Biotechnology
            Genome Institute of Singapore
            1 Science Park Road, The Capricorn #05-01, Singapore 117528
            Tel: +65 6827 5200
            Fax: +65 6827 5201
            Email: gisry@nus.edu.sg
            GIS Clone ID: ZF001-P00031-PP D16
            PCR PRimers
            FORWARD: M13
            BACKWARD: M13
            Plate: ZF001-P00031-DPE-F-D
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                     Location/Qualifiers
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                     /dev stage="7 Different embryonic Stages (From just
                     fertilized Embryos to 72 hours just hatched baby fish)"
                     /lab host="DH10B"
                     /clone lib="GISZF001"
                     /note="Vector: pDNR-LIB; Site 1: Sfi A (GGCCATTACGGCC);
                     Site 2: Sfi B (GGCCGCCTCGGCC); Priming method: Sfi-(dT)30
                     Primed ; Priming sequence: 5.ATTCTAGA GGCCGAGGCGGCC
                     GACATG(T) 30VN ; Directionally cloned,
                                                                5' cloning site:
                                                     5' linker/adaptor sequence:
                     Sfi A site GGCCATTACGGCC;
                     5.AAGCAGTGGTATCAACGCAGAGTGGCC;
                                                          3' cloning site: Sfi B
                                               3' linker/adaptor sequence: same
                     site GGCCGCCTCGGCC ;
                     as the priming sequence; Average insert size: 2kb; For
                     PCR insert analysis: Use M13 Forward and reverse primers;
                     Library Amplified Recombinants (inserts): 98%; Library
                     complexity: 5x106; Full-length construction (method):
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SMART, a Clontech method; Library constructed by: S. Mathavan, Chia-Lin Wei, and Yijun Ruan Genome Institute of Singapore"

ORIGIN

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28.8%; Score 29.4; DB 14; Length 429;
  Query Match
                         56.8%; Pred. No. 87;
  Best Local Similarity
                                                                           0;
           54; Conservative
                                0; Mismatches
                                                 41;
                                                     Indels
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           6 AGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGT 65
Qу
             216 AGAGGAGAGTGAAGACCTCCAGATTGAAGAAACATTCACAGTCAAACATGAAGAGACTGA 275
Db
          66 TGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGG 100
Qу
                  276 AGAAGCTTTCAGAGTCAAACATGAAGATCCTGAGG 310
Db
RESULT 33
BQ993297/c
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                                                     linear
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           BQ993297
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LOCUS
           QGF28E04.yg.ab1 QG EFGHJ lettuce serriola Lactuca sativa cDNA clone
DEFINITION
           QGF28E04, mRNA sequence.
           BQ993297
ACCESSION
           BQ993297.1 GI:22412832
VERSION
KEYWORDS
           EST.
SOURCE
           Lactuca sativa
  ORGANISM Lactuca sativa
           Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
           Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
           asterids; campanulids; Asterales; Asteraceae; Cichorioideae;
           Cichorieae: Lactuca.
               (bases 1 to 463)
REFERENCE
           Kozik, A., Michelmore, R.W., Knapp, S., Matvienko, M., Rieseberg, L.,
  AUTHORS
           Lin, H., van Damme, M., Lavelle, D., Chevalier, P., Ziegle, J.,
           Ellison, P., Kolkman, J., Slabaugh, M.S., Livingston, K., Zhou, Y.,
           Lai, Z., Church, S., Jackson, L. and Bradford, K.
           Lettuce and Sunflower ESTs from the Compositae Genome Project
  TITLE
           http://compgenomics.ucdavis.edu/
  JOURNAL
           Unpublished (2002)
           Contact: Alexander Kozik [R.W.Michelmore]
COMMENT
           Department of Vegetable Crops, R.W.Michelmore Lab
           University of California at Davis (UCD)
           Asmundson Hall, UCD, Davis, CA 95616, USA
           Tel: 1-(530)-742-1742
            Fax: 1-(530)-752-9659
            Email: akozik@atgc.org [michelmore@vegmail.ucdavis.edu]
            belongs to contig QG CA Contig7941, see http://cgpdb.ucdavis.edu/
            for details.
            Plate: QGF28 row: E column: 04.
                    Location/Qualifiers
FEATURES
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                    /clone="OGF28E04"
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/lab host="E.coli" /clone lib="QG EFGHJ lettuce serriola" /note="Vector: pBRcDNASfiAB; The library was constructed from 10 different sources of RNA from a single genotype. Separate cDNAs were generated using primers that incorporated unique 5' and 3' tags to distinguish each source of RNA. cDNAs were then pooled, size-fractionated, directionally cloned into a custom medium-copy vector and transformations made with four size classes to minimize size bias. Details of each source of RNA and library construction can be obtained at http://cgpdb.ucdavis.edu/ TAG SEQ=Not found" 28.8%; Score 29.4; DB 13; Length 463; 76.6%; Pred. No. 90; Best Local Similarity 0; Mismatches 0; Gaps 0; 11; Indels 36; Conservative 18 TGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64 256 TGACCACCAGAGTTTTGAATTGACCACCGGAGGTGTAGTGGAGCATG 210 GSS 01-SEP-2000 linear CNS02AAN 1021 bp DNATetraodon nigroviridis genome survey sequence PUC-Ori end of clone 251G22 of library G from Tetraodon nigroviridis, genomic survey sequence. AL188312 AL188312.1 GI:7826416 GSS; genome survey sequence. Tetraodon nigroviridis Tetraodon nigroviridis Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes; Tetradontoidea; Tetraodontidae; Tetraodon. Roest Crollius, H., Jaillon, O., Dasilva, C., Bouneau, L., Fisher, C., Bernot, A., Fizames, C., Wincker, P., Brottier, P., Quetier, F., Saurin, W. and Weissenbach, J. Estimate of human gene number provided by genome-wide analysis using Tetraodon nigroviridis DNA sequence Nat. Genet. 25 (2), 235-238 (2000) 20296633 10835645 Roest Crollius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C., Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and Weissenbach, J. Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetraodon nigroviridis Genome Res. 10 (7), 939-949 (2000)

ORIGIN

Qy

Db

RESULT 34 CNS02AAN/c

DEFINITION

ACCESSION

VERSION

KEYWORDS

REFERENCE AUTHORS

TITLE

JOURNAL

MEDLINE PUBMED

AUTHORS

REFERENCE

TITLE

JOURNAL

MEDLINE PUBMED

REFERENCE

20359837

10899143

3 (bases 1 to 1021)

ORGANISM

SOURCE

LOCUS

Query Match

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AUTHORS
           Genoscope.
           Direct Submission
 TITLE
           Submitted (12-APR-2000) Genoscope - Centre National de Sequencage:
 JOURNAL
           BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr
           - Web : www.genoscope.cns.fr)
           This sequence is a single read and was generated as part of a large
COMMENT
           scale clone-end sequencing project of the Tetraodon nigroviridis
           genome. For more information, please take a look at
           http://www.genoscope.cns.fr/Tetraodon.
                    Location/Qualifiers
FEATURES
                    1. .1021
    source
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                    /mol type="genomic DNA"
                    /db xref="taxon:99883"
                    /clone="251G22"
                    /clone lib="G"
                    /note="Genoscope sequence ID : COAG251BD11SP1~end :
                    PUC-Ori"
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                         28.8%; Score 29.4; DB 29;
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                         60.8%; Pred. No. 1.4e+02;
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 Matches
           48; Conservative
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                                                 31;
                                                      Indels
                                                                    Gaps
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Qγ
                                       173 TGTTCTGTCGAGTCTTGGACTTCTGCTTCTGGCTGAAGGTCAGATTGTTGCTGCTGATGG 114
Db
          78 AGGAGAACAAGCTGTCCTG 96
Qy
                  Db
         113 TTCCAAACAAGCTCTCCTG 95
RESULT 35
CA628204
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                                    536 bp
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                                                      linear
LOCUS
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           wlel.pk0005.c7 wlel Triticum aestivum cDNA clone wlel.pk0005.c7 5'
DEFINITION
            end, mRNA sequence.
ACCESSION
           CA628204
VERSION
            CA628204.1 GI:25206500
KEYWORDS
            Triticum aestivum (bread wheat)
SOURCE
           Triticum aestivum
  ORGANISM
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Pooideae; Triticeae; Triticum.
            1 (bases 1 to 536)
REFERENCE
            Tingey, S.V., Powell, W., Wolters, P., Dolan, M., Hainey, C., Yuan, Z.,
  AUTHORS
            Miao, G., Caraher, N. and Hanafey, M.K.
            DuPont Wheat cDNA Sequence
  TITLE
            Unpublished (2002)
  JOURNAL
            Contact: Scott V. Tingey
COMMENT
            Crop Genetics
            E. I. DuPont de Nemours and Company
            1 Innovation Way, P.O. Box 6104, Newark, DE 19714-6104, USA
            Tel: 302-631-2602
```

Fax: 302-631-2607

```
Email: Scott.V.Tingey@USA.dupont.com
            Seq primer: M13.
FEATURES
                    Location/Qualifiers
                    1. .536
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                     /organism="Triticum aestivum"
                     /mol type="mRNA"
                     /db xref="taxon:4565"
                     /clone="wle1.pk0005.c7"
                     /tissue type="leaf"
                     /clone lib="wle1"
                     /note="Vector: pBluescript SK+; Site 1: EcoRI; Site 2:
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                    etiolated seedling"
ORIGIN
                         28.6%; Score 29.2; DB 14; Length 536;
 Query Match
 Best Local Similarity 74.0%; Pred. No. 1.1e+02;
           37; Conservative
                                0; Mismatches
                                                  13; Indels
                                                               0; Gaps
                                                                             0;
           2 TGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGT 51
Qу
                    Db
         447 TGGTGTCTGAGCTCTCAAACCTCCAGAGTGATGGTCTTGCCGGTGAAGGT 496
RESULT 36
B0743419/c
LOCUS
            B0743419
                                    674 bp
                                              mRNA
                                                       linear
                                                                EST 17-JUL-2002
           WHE4103 G06 N11ZS Wheat salt-stressed root cDNA library Triticum
DEFINITION
            aestivum cDNA clone WHE4103 G06 N11, mRNA sequence.
ACCESSION
           B0743419
VERSION
           BO743419.1 GI:21890206
KEYWORDS
           EST.
            Triticum aestivum (bread wheat)
SOURCE
 ORGANISM
           Triticum aestivum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Pooideae; Triticeae; Triticum.
               (bases 1 to 674)
REFERENCE
 AUTHORS
           Anderson, O.D., Akhunov, E., Chao, S., Crossman, C., Deal, K.,
            Dvorak, J., Lazo, G.R., Pham, J., Rausch, C.J., Wilson, C. and Woo, J.
 TITLE
            The structure and function of the expressed portion of the wheat
            genomes - Salt-stressed root cDNA library
  JOURNAL
            Unpublished (2002)
            Contact: Olin Anderson
COMMENT
            US Department of Agriculture, Agriculture Research Service, Pacific
            West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
            Tel: 5105595773
            Fax: 5105595818
            Email: oandersn@pw.usda.gov
            Sequences have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seg primer: SK primer.
                    Location/Qualifiers
FEATURES
                     1. .674
     source
                     /organism="Triticum aestivum"
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/mol type="mRNA"

/clone="WHE4103 G06 N11" /tissue type="Roots" /dev stage="Full tillering" /lab host="E. coli SOLR" /clone lib="Wheat salt-stressed root cDNA library" /note="Vector: Lambda Uni-ZAP XR, excised phagemid pBluescript SK(-); Site 1: EcoRI; Site 2: XhoI; Hydroponic plants grown to full tillering stage were treated with 150 mM NaCl for either 12 hours or 7 days. Root tissues of the plants subjected to both types of treatment were collected separately at University of California, Davis (E. Akhunov and K. Deal in J. Dvorak's Lab). Total RNA was prepared separately from the two samples (12h and 7day treatments), and equal amount of RNA was then pooled. PolyA RNA was purified from the pooled RNA, a cDNA library was made, and the cDNA clones were in vivo excised to give pBluescript SK(-) phagemids in J. Dvorak's lab (E. Akhunov, J. Dvorak) at the University of California, Davis. Colony plating, plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors)."

ORIGIN

Query Match Best Local Similarity								Lengt	h 674	1;		
Matches			_					Indel	s	0;	Gaps	0;
Qу	1 (CTGGTAGG	TGAG	SATCTCT				CCACTG				60
Db	99 (CTGGTGCC	CGCA	GTCCTG	 							40
QУ	61 A	ACTGTTGT	CACT	TTCCGA	 SAACZ	 TCCT	GGA 9	8				
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/cultivar="Chinese Spring"
/db xref="taxon:4565"

RESULT 37 BH489764

LOCUS BH489764 362 bp DNA linear GSS 13-DEC-2001

DEFINITION BOHQC87TR BOHQ Brassica oleracea genomic clone BOHQC87, genomic

survey sequence.

ACCESSION BH489764

VERSION BH489764.1 GI:17697868

KEYWORDS GSS.

SOURCE Brassica oleracea
ORGANISM Brassica oleracea

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Brassica.

REFERENCE 1 (bases 1 to 362)

AUTHORS Town, C.D., Van Aken, S., Utterback, T., Koo, H. and Fraser, C.M.

TITLE Whole genome shotgun sequencing of Brassica oleracea

JOURNAL Unpublished (2001)
COMMENT Other GSSs: BOHQC87TF

Contact: Chris Town

TIGR

```
9712 Medical Center Drive, Rockville, MD 20850, USA.
           Tel: 301-838-3523
           Fax: 301-838-0208
           Email: cdtown@tigr.org
           DNA is from a doubled haploid provided by Tom Osborn.
           Seq primer: TR
           Class: sheared ends.
FEATURES
                    Location/Qualifiers
                    1. .362
     source
                    /organism="Brassica oleracea"
                    /mol_type="genomic DNA"
                    /strain="TO1000DH3"
                    /db xref="taxon:3712"
                    /clone="BOHQC87"
                    /clone lib="BOHQ"
                    /note="Vector: pHOS1; Site 1: BstXI; 2-3 kb sheared
                    genomic DNA inserted into pHOS1 using BstXI linkers"
ORIGIN
  Query Match
                         28.4%; Score 29; DB 28; Length 362;
  Best Local Similarity
                         58.8%; Pred. No. 1.1e+02;
           50; Conservative
                                0; Mismatches
                                                 35; Indels
                                                                   Gaps
                                                                            0;
          14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
Qγ
                                          1 | 1 | 11|||| 1 | 11|| 1 | 1
              168 TTTCTAAGCATCAAGGTGTTGCATCTGTTATTTTTGATGAAGTGGATACTGGTGTAAGTG 227
Db
          74 TCCGAGGAGAACAAGCTGTCCTGGA 98
Qу
               228 GCCGGGTCGCACAGGCTATTGCGGA 252
Db
RESULT 38
BU046816/c
                                                      linear
                                                               EST 26-AUG-2002
LOCUS
            BU046816
                                     436 bp
                                              mRNA
           PP LEa0027M13f Peach developing fruit mesocarp Prunus persica cDNA
DEFINITION
            clone PP LEa0027M13f, mRNA sequence.
            BU046816
ACCESSION
           BU046816.1 GI:22486893
VERSION
KEYWORDS
            EST.
SOURCE
            Prunus persica (peach)
           Prunus persica
  ORGANISM
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids I; Rosales; Rosaceae; Amygdaloideae; Prunus.
            1 (bases 1 to 436)
REFERENCE
            Callahan, A., Palmer, M., Main, D., Wing, R. and Abbott, A.
  AUTHORS
            Peach Model Genome for Rosaceae
  TITLE
            Unpublished (2002)
  JOURNAL
            Contact: Abbott, A.
COMMENT
            Dept of Genetics and Biochemistry
            Clemson University
            122 Long Hall, Clemson University, Clemson, SC 29634, USA
            Tel: 864 656 3060
            Fax: 864 656 6879
            Email: aalbert@clemson.edu
            Total High Quality bases = 328
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Seq primer: TAATACGACTCACTATAGGG
            High quality sequence stop: 436.
FEATURES
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                    1. .436
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                    /mol type="mRNA"
                    /cultivar="Loring"
                    /db xref="taxon:3760"
                    /clone="PP LEa0027M13f"
                    /tissue type="Mesocarp"
                    /lab host="E. coli"
                    /clone lib="Peach developing fruit mesocarp"
                    /note="Vector: pBluescript II SK(-); Site_1: EcoRI;
                    Site 2: XhoI; authority=Prunus persica L. Batsh; The
                    sequence has been trimmed to remove vector sequence and
                    contains a minimum of 100 bases of phred value 20 or
                    above. For more details on library preparation and
                    sequence analysis go to
                    http://www.genome.clemson.edu/projects/peach. To order
                    this clone go to http://www.genome.clemson.edu/orders"
ORIGIN
  Query Match
                         28.4%; Score 29; DB 13; Length 436;
  Best Local Similarity
                         58.8%; Pred. No. 1.2e+02;
 Matches
          50; Conservative
                                0; Mismatches
                                                 35;
                                                     Indels
                                                                0; Gaps
                                                                            0;
Qу
           11 AGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCA 70
              1 11 11 111 1111 11
                                       315 ACATTTCCGACGACCAGATTGGCCTTCTTCCCACAGTAGATGAACTGGCCAGTGTAGATA 256
Db
Qy
          71 CTTTCCGAGGAGAACAAGCTGTCCT 95
              255 CCCTCAGCGGCGACGAAGAGCTCGT 231
Db
RESULT 39
BU042469/c
LOCUS
                                    614 bp
                                              mRNA
                                                      linear
                                                               EST 26-AUG-2002
            BU042469
DEFINITION
          PP LEa0012L15f Peach developing fruit mesocarp Prunus persica cDNA
            clone PP LEa0012L15f, mRNA sequence.
ACCESSION
            BU042469
            BU042469.1 GI:22482546
VERSION
KEYWORDS
            EST.
            Prunus persica (peach)
SOURCE
            Prunus persica
  ORGANISM
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids I; Rosales; Rosaceae; Amygdaloideae; Prunus.
               (bases 1 to 614)
REFERENCE
            Callahan, A., Palmer, M., Main, D., Wing, R. and Abbott, A.
  AUTHORS
            Peach Model Genome for Rosaceae
  TITLE
            Unpublished (2002)
  JOURNAL
COMMENT
            Contact: Abbott, A.
            Dept of Genetics and Biochemistry
            Clemson University
```

122 Long Hall, Clemson University, Clemson, SC 29634, USA

Tel: 864 656 3060

```
Email: aalbert@clemson.edu
           Total High Quality bases = 498
           Seg primer: TAATACGACTCACTATAGGG
           High quality sequence stop: 614.
FEATURES
                    Location/Qualifiers
                    1. .614
    source
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                    /mol type="mRNA"
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                    /db xref="taxon:3760"
                    /clone="PP LEa0012L15f"
                    /tissue type="Mesocarp"
                    /lab host="E. coli"
                    /clone lib="Peach developing fruit mesocarp"
                    /note="Vector: pBluescript II SK(-); Site 1: EcoRI;
                    Site 2: XhoI; authority=Prunus persica L. Batsh; The
                    sequence has been trimmed to remove vector sequence and
                    contains a minimum of 100 bases of phred value 20 or
                    above. For more details on library preparation and
                    sequence analysis go to
                    http://www.genome.clemson.edu/projects/peach. To order
                    this clone go to http://www.genome.clemson.edu/orders"
ORIGIN
  Query Match
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                                 Score 29; DB 13; Length 614;
                                Pred. No. 1.4e+02;
  Best Local Similarity
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          50; Conservative
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                                                     Indels
                                                                0;
                                                                   Gaps
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              330 ACATTTCCGACGACCAGATTGGCTTTCTTCCCACAGTAGATGAACTGGCCAGTGTAGATA 271
Db
Qy
          71 CTTTCCGAGGAGAACAAGCTGTCCT 95
              270 CCCTCAGCGGCGACGAAGAGCTCGT 246
Db
RESULT 40
BU046581/c
LOCUS
           BU046581
                                    630 bp
                                              mRNA
                                                      linear
                                                               EST 26-AUG-2002
           PP LEa0026M12f Peach developing fruit mesocarp Prunus persica cDNA
DEFINITION
           clone PP LEa0026M12f, mRNA sequence.
ACCESSION
           BU046581
           BU046581.1 GI:22486658
VERSION
           EST.
KEYWORDS
SOURCE
           Prunus persica (peach)
           Prunus persica
  ORGANISM
           Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
           Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids I; Rosales; Rosaceae; Amygdaloideae; Prunus.
               (bases 1 to 630)
REFERENCE
           1
           Callahan, A., Palmer, M., Main, D., Wing, R. and Abbott, A.
  AUTHORS
            Peach Model Genome for Rosaceae
  TITLE
  JOURNAL
           Unpublished (2002)
COMMENT
           Contact: Abbott, A.
            Dept of Genetics and Biochemistry
```

Fax: 864 656 6879

```
Clemson University
           122 Long Hall, Clemson University, Clemson, SC 29634, USA
           Tel: 864 656 3060
           Fax: 864 656 6879
           Email: aalbert@clemson.edu
           Total High Quality bases = 523
            Seg primer: TAATACGACTCACTATAGGG
           High quality sequence stop: 630.
                    Location/Qualifiers
FEATURES
                    1. .630
    source
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                    /clone="PP LEa0026M12f"
                    /tissue type="Mesocarp"
                    /lab host="E. coli"
                    /clone lib="Peach developing fruit mesocarp"
                     /note="Vector: pBluescript II SK(-); Site 1: EcoRI;
                    Site 2: XhoI; authority=Prunus persica L. Batsh; The
                    sequence has been trimmed to remove vector sequence and
                    contains a minimum of 100 bases of phred value 20 or
                    above. For more details on library preparation and
                    sequence analysis go to
                    http://www.genome.clemson.edu/projects/peach. To order
                    this clone go to http://www.genome.clemson.edu/orders"
ORIGIN
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Qy
                                       1 11 11 111 1111 11
                                                               332 ACATTTCCGACGACCAGATTGGCCTTCTTCCCACAGTAGATGAACTGGCCAGTGTAGATA 273
Db
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Qу
              111
          272 CCCTCAGCGGCGACGAAGAGCTCGT 248
Db
RESULT 41
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                                                               EST 26-AUG-2002
LOCUS
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            PP LEa0018018f Peach developing fruit mesocarp Prunus persica cDNA
DEFINITION
            clone PP LEa0018018f, mRNA sequence.
            BU044321
ACCESSION
            BU044321.1 GI:22484398
VERSION
KEYWORDS
            EST.
            Prunus persica (peach)
SOURCE
  ORGANISM
            Prunus persica
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids I; Rosales; Rosaceae; Amygdaloideae; Prunus.
               (bases 1 to 635)
REFERENCE
            Callahan, A., Palmer, M., Main, D., Wing, R. and Abbott, A.
  AUTHORS
            Peach Model Genome for Rosaceae
  TITLE
```

```
Unpublished (2002)
COMMENT
            Contact: Abbott, A.
            Dept of Genetics and Biochemistry
            Clemson University
            122 Long Hall, Clemson University, Clemson, SC 29634, USA
            Tel: 864 656 3060
            Fax: 864 656 6879
            Email: aalbert@clemson.edu
            Total High Quality bases = 522
            Seq primer: TAATACGACTCACTATAGGG
            High quality sequence stop: 635.
                     Location/Qualifiers
FEATURES
                     1. .635
     source
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                     /lab host="E. coli"
                     /clone lib="Peach developing fruit mesocarp"
                     /note="Vector: pBluescript II SK(-); Site 1: EcoRI;
                     Site 2: XhoI; authority=Prunus persica L. Batsh; The
                     sequence has been trimmed to remove vector sequence and
                     contains a minimum of 100 bases of phred value 20 or
                     above. For more details on library preparation and
                     sequence analysis go to
                     http://www.genome.clemson.edu/projects/peach. To order
                     this clone go to http://www.genome.clemson.edu/orders"
ORIGIN
                          28.4%; Score 29; DB 13; Length 635;
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                          58.8%; Pred. No. 1.4e+02;
  Best Local Similarity
            50; Conservative
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 Matches
                                 0; Mismatches
                                                 35; Indels
                                                                 0; Gaps
           11 AGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCA 70
Qу
                                                                I \cap I \cap I
                                        11 1111 1111 1111 1
              1 11 11 111 1111 11
          315 ACATTTCCGACGACCAGATTGGCCTTCTTCCCACAGTAGATGAACTGGCCAGTGTAGATA 256
Db
           71 CTTTCCGAGGAGAACAAGCTGTCCT 95
Qу
              1 11 1 11 11 111
                                  255 CCCTCAGCGGCGACGAAGAGCTCGT 231
Db
RESULT 42
BH109216/c
                                                                GSS 19-JUL-2001
                                     735 bp
                                                       linear
            BH109216
                                               DNA
LOCUS
            RPCI-24-340C23.TJ RPCI-24 Mus musculus genomic clone
DEFINITION
            RPCI-24-340C23, genomic survey sequence.
            BH109216
ACCESSION
            BH109216.1 GI:14942075
VERSION
KEYWORDS
            GSS.
            Mus musculus (house mouse)
SOURCE
  ORGANISM Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
            1 (bases 1 to 735)
REFERENCE
```

JOURNAL

```
Tsegaye, G., Geer, K., Krol, M., Shvartsbeyn, A., Gebregeorgis, E.,
           Russell, D., de Jong, P. and Fraser, C.M.
 TITLE
           Mouse BAC End Sequences from Library RPCI-24
           Unpublished (1999)
 JOURNAL
           Other GSSs: RPCI-24-340C23.TV
COMMENT
           Contact: Shaying Zhao
           Department of Eukaryotic Genomics
           The Institute for Genomic Research
           9712 Medical Center Dr., Rockville, MD 20850, USA
           Tel: 301 838 0200
           Fax: 301 838 0208
           Email: szhao@tigr.org
           Clones are derived from the mouse BAC library RPCI-24. For BAC
           library availability, please contact Pieter de Jong
            (pdejong@mail.cho.org). Clones may be purchased from BACPAC
           Resources (http://www.chori.org/bacpac/orderingframe.htm). BAC end
           page: http://ww .tigr.org/tdb/bac ends/mouse/bac end intro.html
           Plate: 340 row: C column: 23
           Seq primer: SP6
           Class: BAC ends.
FEATURES
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                     /mol type="genomic DNA"
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                     /db xref="taxon:10090"
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                     /cell type="Spleen/Brain"
                     /clone lib="RPCI-24"
                     /note="Vector: pTARBAC1; Site 1: BamH1; Site 2: BamH1;
                     RPCI-24 Mouse BAC Library produced by Pieter de Jong. The
                     library was cloned in the pTARBAC1 cloning vector at the
                     BamH1 sites using MboI partially digested male C57BL/6J
                     DNA."
ORIGIN
 Ouery Match
                          28.4%;
                                 Score 29; DB 28; Length 735;
                         63.8%; Pred. No. 1.5e+02;
 Best Local Similarity
                                                 25; Indels
           44; Conservative
                                 0; Mismatches
                                                                 0;
                                                                    Gaps
                                                                             0;
           27 GAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACA 86
Qy
                           111111
                                    1 1
                                                     11111
                                                           666 GATTTCTGGAGCTTCCACTGTCTGTCAAGTTGTGGCACATGTCAGCTCACAAGGAGAACA 607
Db
           87 AGCTGTCCT 95
Qy
              1 111 1 1
          606 AACTGGCTT 598
Db
RESULT 43
AI117880/c
                                                                EST 02-SEP-1998
LOCUS
           AI117880
                                     342 bp
                                               mRNA
                                                       linear
           uc41f02.rl Soares mammary gland NMLMG Mus musculus cDNA clone
DEFINITION
           IMAGE:1400571 5', mRNA sequence.
           AI117880
ACCESSION
```

Zhao, S., Nierman, W., Malek, J., Shatsman, S., Akinret, B., Levins, M.,

AUTHORS

```
VERSION
           AI117880.1 GI:3518204
KEYWORDS
           EST.
SOURCE
           Mus musculus (house mouse)
  ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
               (bases 1 to 342)
 AUTHORS
           Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
           Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
           Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
           Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
           Waterston, R.
 TITLE
           The WashU-HHMI Mouse EST Project
  JOURNAL
           Unpublished (1996)
COMMENT
           Contact: Marra M/Mouse EST Project
           WashU-HHMI Mouse EST Project
           Washington University School of MedicineP
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           MGI:912287
           Seg primer: -28m13 rev2 ET from Amersham
           High quality sequence stop: 297.
FEATURES
                    Location/Oualifiers
                     1. .342
    source
                     /organism="Mus musculus"
                     /mol type="mRNA"
                     /db xref="taxon:10090"
                     /clone="IMAGE:1400571"
                     /sex="female (lactating)"
                     /tissue type="mammary gland"
                     /lab host="DH10B"
                     /clone lib="Soares mammary gland NMLMG"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; 1st strand cDNA was prepared from mammary
                    gland tissue from a lactating female, and was then primed
                    with a Not I - oligo(dT) primer. Double-stranded cDNA was
                    ligated to Eco RI adaptors (Pharmacia), digested with Not
                     I and cloned into the Not I and Eco RI sites of the
                    modified pT7T3 vector. Library is normalized. Library
                     was constructed by Bento Soares and M. Fatima Bonaldo. "
ORIGIN
                          28.2%;
                                 Score 28.8; DB 9;
  Query Match
                                                     Length 342;
  Best Local Similarity
                          58.0%; Pred. No. 1.2e+02;
           51; Conservative
                                0; Mismatches
                                                 37; Indels
  Matches
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                                                                            0;
           14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
Qу
              1 1111
                                                              1.1
Db
          260 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 201
           74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
Qv
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200 TTCGATGAGAGCGATCTGTTCTTGTAGC 173

Db

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RESULT 44
AA177634/c
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LOCUS AA177634 398 bp mRNA linear EST 16-FEB-1997

DEFINITION mt32h12.rl Soares mouse 3NbMS Mus musculus cDNA clone IMAGE:622823

5', mRNA sequence.

ACCESSION AA177634

VERSION

AA177634.1 GI:1758868

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE (bases 1 to 398)

AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,

> Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and

Waterston, R.

TITLE The WashU-HHMI Mouse EST Project

JOURNAL Unpublished (1996)

COMMENT Contact: Marra M/Mouse EST Project

WashU-HHMI Mouse EST Project

Washington University School of MedicineP

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800 Fax: 314 286 1810

Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.

MGI:383647

Seq primer: -28M13 rev2 from Amersham

High quality sequence stop: 371.

FEATURES

Location/Qualifiers

source

1. .398

/organism="Mus musculus"

/mol type="mRNA" /strain="C57BL/6J" /db xref="taxon:10090"

/clone="IMAGE:622823"

/sex="male"

/tissue type="Spleen" /dev stage="4 weeks" /lab host="DH10B"

/clone lib="Soares mouse 3NbMS"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'

3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. RNA provided by Dr. Bertrand Jordan. Library went through three rounds of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

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28.2%; Score 28.8; DB 9; Length 398;
  Query Match
  Best Local Similarity 58.0%; Pred. No. 1.3e+02;
                                0; Mismatches
           51; Conservative
                                               37; Indels
                                                                0; Gaps
                                                                            0;
           14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
QV
              1 111 1 11 111 1 1 1
                                   1 1111
Db
         240 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 181
Qy
           74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
              180 TTCGATGAGAGCGATCTGTTCTTGTAGC 153
Db
RESULT 45
BG550348/c
LOCUS
            BG550348
                                    416 bp
                                              mRNA
                                                      linear
                                                               EST 05-APR-2001
DEFINITION
           947039G04.x2 947 - 2 week shoot from Barkan lab Zea mays cDNA, mRNA
           sequence.
           BG550348
ACCESSION
           BG550348.1 GI:13558993
VERSION .
KEYWORDS
            EST.
SOURCE
            Zea mays
  ORGANISM Zea mays
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
            clade; Panicoideae; Andropogoneae; Zea.
           1 (bases 1 to 416)
REFERENCE
           Walbot, V.
  AUTHORS
           Maize ESTs from various cDNA libraries sequenced at Stanford
  TITLE
            University
           Unpublished (1999)
  JOURNAL
            Contact: Walbot V
COMMENT
            Department of Biological Sciences
            Stanford University
            855 California Ave, Palo Alto, CA 94304, USA
            Tel: 650 723 2227
            Fax: 650 725 8221
            Email: walbot@stanford.edu
            Plate: 947039 row: G column: 04.
                    Location/Qualifiers
FEATURES
                    1. .416
     source
                    /organism="Zea mays"
                    /mol type="mRNA"
                    /cultivar="B73"
                    /db xref="taxon:4577"
                    /tissue_type="leaf and stem, including leaf base"
                    /dev stage="2 week old seedling (3 leaves)"
                    /lab host="XL1-Blue"
                    /clone lib="947 - 2 week shoot from Barkan lab"
                    /note="Organ: shoot; Vector: Lambda ZAP (pBlueScript SK-);
                    Site 1: EcoRI; Site 2: XhoI; Directionally cloned using
                    Stratagene's UniZap XR cDNA cloning kit with the 5' end
                    at the EcoRI site. The library represents 8 x 10e5
                    independent recombinant phage. The plants were greenhouse
                    grown."
```

ORIGIN

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28.2%; Score 28.8; DB 12; Length 416;
 Query Match
 Best Local Similarity 60.0%; Pred. No. 1.3e+02;
 Matches
           48; Conservative
                                0; Mismatches
                                                 32; Indels
                                                                           0;
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qy
                 105 CTATCAAGTGTGTAGTGTCTTCGAGAAGTTTGTAGAGCCTACTGCTGCTGCTGTATAT 46
Db
          61 ACTGTTGTCACTTTCCGAGG 80
Qу
             1111 1 11 111 11 111
          45 ACTGATATCGCTTGCCAAGG 26
Db
RESULT 46
BQ557757
                                    510 bp
                                              mRNA
                                                      linear
                                                              EST 20-JUN-2002
LOCUS
           BQ557757
           H4048B01-3 NIA Mouse 7.4K cDNA Clone Set Mus musculus cDNA clone
DEFINITION
           H4048B01 3', mRNA sequence.
           BQ557757
ACCESSION
           BQ557757.1 GI:21458642
VERSION
KEYWORDS
SOURCE
           Mus musculus (house mouse)
  ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
           1 (bases 1 to 510)
           VanBuren, V., Piao, Y., Dudekula, D.B., Qian, Y., Carter, M.G.,
  AUTHORS
           Martin, P.R., Stagg, C.A., Bassey, U., Aiba, K., Hamatani, T.,
           Kargul, G.J., Luo, A.G., Kelso, J., Hide, W. and Ko, M.S.H.
           Assembly, verification, and initial annotation of NIA 7.4K mouse
  TITLE
           cDNA clone set
           Genome Res. 12 (12), 1999-2003 (2002)
  JOURNAL
  MEDLINE
           22354164
   PUBMED
           12466305
           Other ESTs: H4048B01-5
COMMENT
           Contact: Yong Qian
           Laboratory of Genetics
           National Institute on Aging/National Institutes of Health
            333 Cassell Drive, Suite 3000, Baltimore, MD 21224-6820, USA
            Email: cdna@lgsun.grc.nia.nih.gov
           This clone set has been freely distributed to the community. Please
            visit http://lqsun.grc.nia.nih.gov/cDNA/NIA 7 4k.html for details.
            Plate: H4048 row: B column: 01
            Seq primer: -21M13 Forward
            High quality sequence stop: 510
            POLYA=Yes.
                    Location/Qualifiers
FEATURES
                    1. .510
     source
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                    /mol type="mRNA"
                    /strain="C57BL/6"
                    /db xref="niaEST:H4048B01-3"
                    /db xref="taxon:10090"
                    /clone="H4048B01"
                    /sex="mixed"
                    /dev stage="mixed"
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                    /note="Vector: pSPORT1; Site 1: SalI; Site 2: NotI; This
                    clone is among a rearrayed set of 7,407 clones from more
                    than 20 cDNA libraries."
ORIGIN
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                                                     Length 510;
  Query Match
                         58.0%; Pred. No. 1.5e+02;
  Best Local Similarity
                                                               0;
                                                                           0:
           51; Conservative
                                0; Mismatches 37;
                                                     Indels
                                                                   Gaps
          14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
Qу
             11
         150 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 209
Db
          74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
Qу
             210 TTCGATGAGAGCGATCTGTTCTTGTAGC 237
Db
RESULT 47
BX514645/c
LOCUS
           BX514645
                                    524 bp
                                              mRNA
                                                      linear
                                                              EST 25-JUN-2003
           BX514645 Soares mouse 3NbMS Mus musculus cDNA clone IMAGp952C2329;
DEFINITION
           IMAGE: 622823, mRNA sequence.
ACCESSION
           BX514645
VERSION
           BX514645.1 GI:32244604
KEYWORDS
           EST.
           Mus musculus (house mouse)
SOURCE
  ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
           1 (bases 1 to 524)
REFERENCE
           Heil, O., Ebert, L., Neubert, P., Peters, M., Radelof, U., Schneider, D.
  AUTHORS
            and Korn, B.
           Mouse UnigeneSet - RZPD2
  TITLE
           Unpublished (2003)
  JOURNAL
COMMENT
           Contact: Ina Rolfs
            RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
            Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
           RZPD; IMAGp952C2329.
           RZPDLIB; I.M.A.G.E. cDNA Clone Collection;
           Mouse UniqueeSet - RZPD2 (RZPDLIB No.981)
           http://www.rzpd.de/CloneCards/cgi-
           bin/showLib.pl.cgi/response?libNo=981 Contact: Ina Rolfs
            RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
            Heubnerweg 6, D-14059 Berlin, Germany
            Tel: +49 30 32639 101
            Fax: +49 30 32639 111
            www.rzpd.de
            This clone is available royalty-free from RZPD;
            contact RZPD (clone@rzpd.de) for further information. Seg primer:
            T7, Primer sequence: TAATACGACTCACTATAGGG.
                    Location/Qualifiers
FEATURES
                    1. .524
     source
                    /organism="Mus musculus"
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/mol type="mRNA"

/lab host="DH10B"

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                 /db xref="taxon:10090"
                 /clone="IMAGp952C2329; IMAGE:622823"
                 /sex="male"
                 /tissue_type="Spleen"
                 /dev stage="4 weeks"
                 /lab host="DH10B"
                 /clone lib="Soares mouse 3NbMS"
                 /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                 polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
                 was primed with a Not I - oligo(dT) primer [5'
                 3']; double-stranded cDNA was ligated to Eco RI adaptors
                  (Pharmacia), digested with Not I and cloned into the Not I
                 and Eco RI sites of the modified pT7T3 vector. RNA
                 provided by Dr. Bertrand Jordan. Library went through
                 three rounds of normalization, and was constructed by
                 Bento Soares and M.Fatima Bonaldo."
                      28.2%; Score 28.8; DB 13; Length 524;
                      58.0%; Pred. No. 1.5e+02;
Best Local Similarity
         51; Conservative
                            0; Mismatches
                                             37;
                                                  Indels
                                                               Gaps
                                                                      0;
        14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
           \Box
       238 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 179
        74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
           178 TTCGATGAGAGCGATCTGTTCTTGTAGC 151
         BX520764
                                 536 bp
                                          mRNA
                                                  linear
                                                          EST 27-JUN-2003
        BX520764 Soares mammary gland NMLMG Mus musculus cDNA clone
         IMAGp998K043537 ; IMAGE:1400571, mRNA sequence.
         BX520764
         BX520764.1 GI:32301442
         Mus musculus (house mouse)
         Mus musculus
         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
         Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
         1 (bases 1 to 536)
         Heil, O., Ebert, L., Neubert, P., Peters, M., Radelof, U., Schneider, D.
         and Korn, B.
         Mouse UnigeneSet - RZPD2
         Unpublished (2003)
         Contact: Ina Rolfs
         RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
         Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
         RZPD; IMAGp998K043537.
         RZPDLIB; I.M.A.G.E. cDNA Clone Collection;
         Mouse UnigeneSet - RZPD2 (RZPDLIB No.981)
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ORIGIN

Qy

Db

Qу

Db

RESULT 48 BX520764/c

DEFINITION

ACCESSION

EST

http://www.rzpd.de/CloneCards/cgi-

VERSION

SOURCE

KEYWORDS

REFERENCE

TITLE

AUTHORS

JOURNAL COMMENT

ORGANISM

LOCUS

Query Match

Matches

```
bin/showLib.pl.cgi/response?libNo=981 Contact: Ina Rolfs
           RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
           Heubnerweg 6, D-14059 Berlin, Germany
           Tel: +49 30 32639 101
           Fax: +49 30 32639 111
           www.rzpd.de
           This clone is available royalty-free from RZPD;
           contact RZPD (clone@rzpd.de) for further information. Seq primer:
           T7, Primer sequence: TAATACGACTCACTATAGGG.
                    Location/Qualifiers
FEATURES
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                    /mol type="mRNA"
                    /db xref="taxon:10090"
                    /clone="IMAGp998K043537 ; IMAGE:1400571"
                    /sex="female (lactating)"
                    /tissue type="mammary gland"
                    /lab host="DH10B"
                    /clone lib="Soares mammary gland NMLMG"
                    /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; 1st strand cDNA was prepared from mammary
                    gland tissue from a lactating female, and was then primed
                    with a Not I - oligo(dT) primer. Double-stranded cDNA was
                    ligated to Eco RI adaptors (Pharmacia), digested with Not
                    I and cloned into the Not I and Eco RI sites of the
                    modified pT7T3 vector. Library is normalized. Library
                    was constructed by Bento Soares and M. Fatima Bonaldo. "
ORIGIN
                         28.2%; Score 28.8; DB 13; Length 536;
  Query Match
                         58.0%; Pred. No. 1.5e+02;
  Best Local Similarity
                                                                   Gaps
                                                                           0;
                                0; Mismatches
                                                 37; Indels
 Matches
           51; Conservative
          14 TCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTT 73
Qу
                                                             \mathbf{H}
                                                                   271 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 212
Db
          74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
Qу
              Db
         211 TTCGATGAGAGCGATCTGTTCTTGTAGC 184
RESULT 49
AI591944/c
                                    598 bp
                                              mRNA
                                                      linear
                                                               EST 15-MAR-2000
           AI591944
LOCUS
           mt32h12.yl Soares mouse 3NbMS Mus musculus cDNA clone IMAGE:622823
DEFINITION
           5', mRNA sequence.
           AI591944
ACCESSION
           AI591944.1 GI:4600992
VERSION
KEYWORDS
           Mus musculus (house mouse)
SOURCE
  ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
            1 (bases 1 to 598)
           Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
  AUTHORS
           Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y.,
```

```
Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R.,
           Waterston, R. and Wilson, R.
 TITLE
           The WashU-NCI Mouse EST Project 1999
           Unpublished (1999)
 JOURNAL
           Contact: Marra M/WashU-NCI Mouse EST Project 1999
COMMENT
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           This read is a RESEQUENCE of a previously sequenced mouse clone
           This read has been verified (found to hit its original self in the
           correct orientation)
           Putative full length read
           vector to vector length is 915
           MGI:383647
           Seq primer: -40RP from Gibco
           High quality sequence stop: 460
           POLYA=No.
FEATURES
                    Location/Qualifiers
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                    /mol type="mRNA"
                    /strain="C57BL/6J"
                    /db xref="taxon:10090"
                    /clone="IMAGE: 622823"
                    /sex="male"
                    /tissue type="Spleen"
                    /dev stage="4 weeks"
                    /lab host="DH10B"
                    /clone lib="Soares mouse 3NbMS"
                    /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
                    was primed with a Not I - oligo(dT) primer [5'
                    3']; double-stranded cDNA was ligated to Eco RI adaptors
                    (Pharmacia), digested with Not I and cloned into the Not I
                    and Eco RI sites of the modified pT7T3 vector. RNA
                    provided by Dr. Bertrand Jordan. Library went through
                    three rounds of normalization, and was constructed by
                    Bento Soares and M. Fatima Bonaldo."
ORIGIN
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                                                   Length 598;
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                         58.0%;
                                Pred. No. 1.6e+02;
  Best Local Similarity
                               0; Mismatches
                                                37; Indels
           51; Conservative
  Matches
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Qy
             1 1111
         240 TTTCTCAACTATAGAATCTAGTTGTGAAGACTTTTCATTAAGTTGCTCTTGAGAACACTT 181
Db
Qу
          74 TCCGAGGAGAACAAGCTGTCCTGGAGGC 101
             180 TTCGATGAGAGCGATCTGTTCTTGTAGC 153
Db
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Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R.,

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RESULT 50
DR36H15S/c
                                                               GSS 22-NOV-2002
                                    654 bp
                                              DNA
                                                      linear
LOCUS
           DR36H15S
           Danio rerio genomic clone DKEY-36H15, genomic survey sequence.
DEFINITION
ACCESSION
           AL987137
           AL987137.1 GI:25180574
VERSION
KEYWORDS
            GSS.
SOURCE
            Danio rerio (zebrafish)
           Danio rerio
  ORGANISM
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
            Cypriniformes; Cyprinidae; Danio.
            1 (bases 1 to 654)
REFERENCE
           Humphray, S.J., Huckle, E. and Hunt, S.E.
  AUTHORS
            Direct Submission
  TITLE
            Submitted (14-NOV-2002) The Sanger Institute, Wellcome Trust Genome
  JOURNAL
            Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail contact:
            humquery@sanger.ac.uk Unpublished
            This sequence was generated from the SP6 end of BAC 36H15. 36H15 is
COMMENT
            part of the Daniokey BAC Library created by R. Plasterk and N.V.
            Keygene.
            Further details: http://www.sanger.ac.uk/Projects/D_rerio/.
                     Location/Qualifiers
FEATURES
                     1. .654
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                     /clone="DKEY-36H15"
                     /tissue type="Testis"
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Qy
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           69 CACTTTCCGAGGAGAACAAGC 89
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Search completed: April 29, 2004, 18:39:20 Job time: 338.622 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model-

Run on: April 29, 2004, 14:53:09; Search time 435.147 Seconds

(without alignments)

10159.758 Million cell updates/sec

Title: US-09-989-981A-9_COPY_3_104

Perfect score: 102

Sequence: 1 ctggtaggtgagatctctga.....aacaagctgtcctggaggcc 102

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 21671516995 residues

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

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5: gb ov:*

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7: gb ph:*

8: gb pl:*

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10: gb_ro:*

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12. -----

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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С	2	102	100.0	2019	6	AX685731		AX685731 Sequence
С	3	102	100.0	2284	10	AY196216		AY196216 Mus muscu
С	4	102	100.0	3674	10	AF324495		AF324495 Mus muscu
	5	102	100.0	6043	6	AX685737		AX685737 Sequence
С	6	100.4	98.4	2285	10	AY196215		AY196215 Mus muscu
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С	19	84.4	82.7	185045	2	AC146466		AC146466 Callithri
	20	82.8	81.2	202533	2	AC146464		AC146464 Saimiri s
С	21	82.8	81.2	207760	2	AC146286		AC146286 Callicebu
С	22	52.4	51.4	135280	2	AC146282		AC146282 Takifugu
	23	35.8	35.1	169570	5	AL928999		AL928999 Zebrafish
С	24	34.2	33.5	190952	5	BX004832		BX004832 Zebrafish
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	29	32.2	31.6	261	9	MFUSCFTR11		AF162357 Macaca fu
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	31	32.2	31.6	261	9	PHACFTR11		AF162411 Papio ham
	32	32.2	31.6	261	9	RMCFTR11		AF016934 Macaca mu
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47	32.2	31.6	4443	6	AR240927	AR240927 Sequence
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LOCUS
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DEFINITION Mus musculus sterolin 2 (Abcg8) gene, exon 2.
ACCESSION
            AF351800
            AF351800.1 GI:18996438
VERSION
KEYWORDS
            2 of 13
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SOURCE
            Mus musculus (house mouse)
            Mus musculus
  ORGANISM
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
            1 (bases 1 to 1017)
REFERENCE
            Lu, K., Lee, M.-H., Yu, H., Zhou, Y., Sandell, S.A., Salen, G. and
  AUTHORS
            Patel, S.B.
  TITLE
            Molecular cloning, genomic organization, genetic variations, and
            characterization of murine sterolin genes Abcg5 and Abcg8
            J. Lipid Res. 43 (4), 565-578 (2002)
  JOURNAL
            21904563
  MEDLINE
   PUBMED
            11907139
            2 (bases 1 to 1017)
REFERENCE
  AUTHORS
            Lu, K., Zhou, Y., Lee, M.-H. and Patel, S.B.
            Direct Submission
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            Submitted (21-FEB-2001) Division of Endocrinology, Diabetes and
  JOURNAL
            Medical Genetics, Medical University of South Carolina, 114 Doughty
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RESULT 2 AX685731/c

LOCUS AX685731 2019 bp DNA linear PAT 29-MAR-2003

DEFINITION Sequence 3 from Patent WO02081691.

ACCESSION AX685731

VERSION AX685731.1 GI:29371740

KEYWORDS

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1

AUTHORS Hobbs, H.H., Shan, B., Barnes, R. and Tian, H.

TITLE Abcq5 and abcq8: compositions and methods of use

JOURNAL Patent: WO 02081691-A 3 17-OCT-2002;

Tularik Inc. (US); BOARD OF REGENTS UNIVERSITY OF TEXAS SYSTEM

(US)

FEATURES Location/Qualifiers

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DEFINITION
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ACCESSION
           AY196216.1 GI:31322261
VERSION
KEYWORDS
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           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
           1 (bases 1 to 2284)
 AUTHORS
           Wittenburg, H., Lyons, M.A., Li, R., Churchill, G.A., Carey, M.C. and
           Paigen, B.
 TITLE
           Primary Roles of FXR and ABCG5/ABCG8 in Cholesterol Gallstone
           Susceptibility: Evidence from a Cross of PERA/Ei and I/Ln Inbred
           Mice
 JOURNAL
           Unpublished
           2 (bases 1 to 2284)
REFERENCE
 AUTHORS
           Lyons, M.A., Wittenburg, H., Walsh, K.A., Carey, M.C. and Paigen, B.
 TITLE
           Direct Submission
  JOURNAL
           Submitted (12-DEC-2002) The Jackson Laboratory, 600 Main Street,
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DEFINITION Mus musculus sterolin-2 (Abcg8) mRNA, complete cds.

ACCESSION AF324495

VERSION AF324495.1 GI:15088541

KEYWORDS

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 3674)

AUTHORS Lu, K., Lee, M.H., Hazard, S., Brooks-Wilson, A., Hidaka, H., Kojima, H., Ose, L., Stalenhoef, A.F., Mietinnen, T., Bjorkhem, I., Bruckert, E., Pandya, A., Brewer, H.B. Jr., Salen, G., Dean, M., Srivastava, A. and Patel, S.B.

TITLE Two genes that map to the STSL locus cause sitosterolemia: genomic structure and spectrum of mutations involving sterolin-1 and sterolin-2, encoded by ABCG5 and ABCG8, respectively

JOURNAL Am. J. Hum. Genet. 69 (2), 278-290 (2001)

MEDLINE 21344600 PUBMED 11452359

REFERENCE 2 (bases 1 to 3674)

AUTHORS Lu, K., Lee, M.-H. and Patel, S.B.

TITLE Direct Submission

JOURNAL Submitted (29-NOV-2000) Division of Endocrinology, Diabetes and Medical Genetics, Medical University of South Carolina, 114 Doughty

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ACCESSION
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VERSION
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           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
 AUTHORS
           Hobbs, H.H., Shan, B., Barnes, R. and Tian, H.
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Street, STB541, Charleston, SC 29403, USA Location/Qualifiers

FEATURES

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Abcq5 and abcq8: compositions and methods of use
  TITLE
  JOURNAL
           Patent: WO 02081691-A 9 17-OCT-2002;
           Tularik Inc. (US); BOARD OF REGENTS UNIVERSITY OF TEXAS SYSTEM
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ACCESSION
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           AY196215.1 GI:31322259
VERSION
KEYWORDS
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SOURCE
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REFERENCE
           1 (bases 1 to 2285)
 AUTHORS
           Wittenburg, H., Lyons, M.A., Li, R., Churchill, G.A., Carey, M.C. and
 TITLE
           Primary Roles of FXR and ABCG5/ABCG8 in Cholesterol Gallstone
           Susceptibility: Evidence from a Cross of PERA/Ei and I/Ln Inbred
           Mice
 JOURNAL
           Unpublished
REFERENCE
           2 (bases 1 to 2285)
 AUTHORS
           Lyons, M.A., Wittenburg, H., Walsh, K.A., Carey, M.C. and Paigen, B.
           Direct Submission
 TITLE
 JOURNAL
           Submitted (12-DEC-2002) The Jackson Laboratory, 600 Main Street,
           Bar Harbor, ME 04609, USA
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                                                              ROD 26-AUG-2002
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DEFINITION
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ACCESSION
VERSION
           AF351785.2 GI:22477145
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REFERENCE
              (bases 1 to 4829)
           Lu, K., Lee, M.H., Hazard, S., Brooks-Wilson, A., Hidaka, H., Kojima, H.,
  AUTHORS
           Ose, L., Stalenhoef, A.F., Mietinnen, T., Bjorkhem, I., Bruckert, E.,
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           Patel, S.B.
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TITLE
            Two genes that map to the STSL locus cause sitosterolemia: genomic
            structure and spectrum of mutations involving sterolin-1 and
            sterolin-2, encoded by ABCG5 and ABCG8, respectively
            Am. J. Hum. Genet. 69 (2), 278-290 (2001)
 JOURNAL
 MEDLINE
            21344600
            11452359
   PUBMED
REFERENCE
            2 (bases 1 to 4829)
 AUTHORS
            Lu, K., Yu, H., Lee, M. and Patel, S.B.
  TITLE
            Molecular cloning, genomic structure, and characterization of novel
            mouse head-to-head tandem ABC transporters
  JOURNAL
            Unpublished
REFERENCE
            3 (bases 1 to 4829)
 AUTHORS
            Lu, K., Lee, M. and Patel, S.B.
 TITLE
            Direct Submission
  JOURNAL
            Submitted (21-FEB-2001) Division of Endocrinology, Diabetes and
            Medical Genetics, Medical University of South Carolina, 114 Doughty
            St, STB 541, Charleston, SC 29407, USA
            4 (bases 1 to 4829)
REFERENCE
  AUTHORS
            Lu, K., Yu, H., Lee, M. and Patel, S.B.
            Direct Submission
  TITLE
            Submitted (26-AUG-2002) Division of Endocrinology, Diabetes and
 JOURNAL
            Medical Genetics, Medical University of South Carolina, 114 Doughty
            St, STB 541, Charleston, SC 29403, USA
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96.0%; Pred. No. 3.8e-21;

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4; Indels

0;

Gaps 0;

Best Local Similarity

97; Conservative

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Qv.
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DEFINITION
           complete cds.
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ACCESSION
           AY145899.1 GI:24935208
VERSION
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             (bases 1 to 40929)
REFERENCE
           Yu, H., Lu, K., Lee, M., Pandit, B. and Patel, s.B.
 AUTHORS
  TITLE
           The rat Abcg5 and Abcg8: characterization, chromosomal assignment
           and genetic variation in sitosterolemic rats
  JOURNAL
           Unpublished
           2 (bases 1 to 40929)
REFERENCE
 AUTHORS
           Yu, H., Lu, K., Lee, M., Pandit, B. and Patel, s.B.
  TITLE
           Direct Submission
           Submitted (29-AUG-2002) Endocrinology, Diabetes and Medical
  JOURNAL
           Genetics, Medical University of South Carolina, 114 Doughty Street,
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4 unordered pieces. ACCESSION AC120701

VERSION AC120701.4 GI:23265381

KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS ENRICHED.

SOURCE Rattus norvegicus (Norway rat)

ORGANISM Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

REFERENCE 1 (bases 1 to 237445)

AUTHORS

Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D.,

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TITLE Direct Submission

JOURNAL Unpublished

REFERENCE (bases 1 to 237445)

AUTHORS Worley, K.C.

TITLE Direct Submission

Submitted (09-MAY-2002) Human Genome Sequencing Center, Department **JOURNAL** of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

REFERENCE 3 (bases 1 to 237445)

AUTHORS Rat Genome Sequencing Consortium.

TITLE Direct Submission

JOURNAL

Submitted (21-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Sep 21, 2002 this sequence version replaced gi:21908396. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequening reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the sequence may extend beyond the ends of the clone and there may be contigs that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only

contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu ----- Project Information

Center project name: GXQV Center clone name: CH230-65H6

----- Summary Statistics

Assembly program: Phrap; version 0.990329 Consensus quality: 209781 bases at least Q40 Consensus quality: 213033 bases at least Q30 Consensus quality: 214997 bases at least Q20

Estimated insert size: 233017; sum-of-contigs estimation Quality coverage: 4x in Q20 bases; sum-of-contigs estimation

COMMENT

^{*} NOTE: Estimated insert size may differ from sequence length

⁽see http://www.hgsc.bcm.tmc.edu/docs/Genbank draft data.html).

^{*} NOTE: This is a 'working draft' sequence. It currently

^{*} consists of 4 contigs. The true order of the pieces

^{*} is not known and their order in this sequence record is

^{*} arbitrary. Gaps between the contigs are represented as

^{*} runs of N, but the exact sizes of the gaps are unknown.

^{*} This record will be updated with the finished sequence

^{*} as soon as it is available and the accession number will

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                       235011: contig of 1045 bp in length
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VERSION
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SOURCE
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REFERENCE
           1 (bases 1 to 312858)
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AUTHORS Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

TITLE Direct Submission JOURNAL Unpublished 2 (bases 1 to 312858) REFERENCE AUTHORS

Worley, K.C.

TITLE Direct Submission

JOURNAL Submitted (24-FEB-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

REFERENCE 3 (bases 1 to 312858) AUTHORS Rat Genome Sequencing Consortium. TITLE Direct Submission Submitted (08-OCT-2002) Human Genome Sequencing Center, Department JOURNAL of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA COMMENT On Sep 23, 2002 this sequence version replaced gi:21738477. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotqun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table. ----- Genome Center Center: Baylor College of Medicine Center code: BCM Web site: http://www.hgsc.bcm.tmc.edu/ Contact: hgsc-help@bcm.tmc.edu ----- Project Information Center project name: GRAX Center clone name: CH230-359E1 ----- Summary Statistics Assembly program: Phrap; version 0.990329 Consensus quality: 241372 bases at least Q40 Consensus quality: 245333 bases at least Q30 Consensus quality: 248022 bases at least Q20 Estimated insert size: 276767; sum-of-contigs estimation Quality coverage: 4x in Q20 bases; sum-of-contigs estimation * NOTE: Estimated insert size may differ from sequence length (see http://www.hqsc.bcm.tmc.edu/docs/Genbank draft data.html) * NOTE: This sequence may represent more than one clone. * NOTE: This is a 'working draft' sequence. It currently * consists of 8 contigs. The true order of the pieces * is not known and their order in this sequence record is * arbitrary. Gaps between the contigs are represented as * runs of N, but the exact sizes of the gaps are unknown. * This record will be updated with the finished sequence * as soon as it is available and the accession number will * be preserved. 1 155105: contig of 155105 bp in length 155106 155205: gap of unknown length 155206 221765: contig of 66560 bp in length 221766 221865: gap of unknown length 221866 290378: contig of 68513 bp in length 290379 290478: gap of unknown length 293724: contig of 3246 bp in length 290479 293725 293824: gap of unknown length 293825 305790: contig of 11966 bp in length 305791 305890: gap of unknown length

307341: contig of 1451 bp in length

305891

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307342
                        307441: gap of unknown length
                        309768: contig of 2327 bp in length
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                        309868: gap of unknown length
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                                                      Indels
                                                                0; Gaps
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Qу
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           61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGC 101
Qу
               88111 GCTGTTGTCACTTTCAGAGGAGAACACGCTGTCCTGGAGGC 88151
Db
RESULT 11
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LOCUS
           AF320294
                                   2022 bp
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                                                      linear
                                                               PRI 13-DEC-2000
DEFINITION
           Homo sapiens ABCG8 (ABCG8) mRNA, complete cds.
ACCESSION
           AF320294
           AF320294.1 GI:11692801
VERSION
KEYWORDS
SOURCE
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           Homo sapiens
  ORGANISM
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
               (bases 1 to 2022)
  AUTHORS
            Berge, K.E., Tian, H., Graf, G.A., Yu, L., Grishin, N.V., Schultz, J.,
            Kwiterovich, P., Shan, B., Barnes, R. and Hobbs, H.H.
  TITLE
            Accumulation of Dietary Cholesterol in Sitosterolemia Caused by
           Mutations in Adjacent ABC Transporters
  JOURNAL
           Science (2001) In press
               (bases 1 to 2022)
REFERENCE
  AUTHORS
            Berge, K.E., Tian, H., Graf, G.A., Yu, L., Grishin, N.V., Schultz, J.,
            Kwiterovich, P., Shan, B., Barnes, R. and Hobbs, H.H.
  TITLE
           Direct Submission
  JOURNAL
           Submitted (09-NOV-2000) Molecular Genetics, University of Texas,
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Dallas, TX 75390-9046, USA
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                                                    Length 2022;
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                                0;
                                   Mismatches
                                                                           0;
                                                     Indels
                                                                   Gaps
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Qу
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Qу
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RESULT 12
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LOCUS
                                   2669 bp
                                             DNA
                                                     linear
                                                              PAT 29-MAR-2003
DEFINITION
           Sequence 7 from Patent WO02081691.
ACCESSION
           AX685735
           AX685735.1 GI:29371744
VERSION
KEYWORDS
SOURCE
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  ORGANISM
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           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
 AUTHORS
           Hobbs, H.H., Shan, B., Barnes, R. and Tian, H.
 TITLE
           Abcq5 and abcq8: compositions and methods of use
```

Southwestern Medical Center at Dallas, 5323 Harry Hines Blvd.,

Patent: WO 02081691-A 7 17-OCT-2002;

Tularik Inc. (US) ; BOARD OF REGENTS UNIVERSITY OF TEXAS SYSTEM

(US)

FEATURES

Location/Qualifiers

source

1. .2669

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CDS

100. .2121

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/codon start=1

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ORIGIN

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Best Local Similarity Pred. No. 8.9e-19; 91.2%;

0; Mismatches Matches 93; Conservative Indels 0; 0; Gaps

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204 GCTGTTGTCACTTTCAGAGGAGAACAATCTATCCTGGAGGCC 163 Db

RESULT 13 AC108476/c

LOCUS AC108476 139342 bp

DEFINITION Homo sapiens BAC clone RP11-1413K20 from 2, complete sequence.

ACCESSION AC108476

AC108476.5 GI:19807988 VERSION

KEYWORDS HTG.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

> Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

DNA

linear

PRI 16-APR-2002

REFERENCE (bases 1 to 139342)

AUTHORS Sulston, J.E. and Waterston, R.

Toward a complete human genome sequence TITLE

Genome Res. 8 (11), 1097-1108 (1998) JOURNAL

MEDLINE 99063792 PUBMED 9847074

REFERENCE 2 (bases 1 to 139342)

AUTHORS Harkins, C., Haakenson, W. and Doebber, A.

TITLE The sequence of Homo sapiens BAC clone RP11-1413K20

JOURNAL Unpublished (2001)

REFERENCE 3 (bases 1 to 139342)

AUTHORS Waterston, R.H.
TITLE Direct Submission

JOURNAL Submitted (27-JAN-2002) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

REFERENCE 4 (bases 1 to 139342)

AUTHORS Waterston, R.H.
TITLE Direct Submission

JOURNAL Submitted (20-FEB-2002) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

REFERENCE 5 (bases 1 to 139342)

AUTHORS Waterston, R.H.
TITLE Direct Submission

JOURNAL Submitted (29-MAR-2002) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

REFERENCE 6 (bases 1 to 139342)

AUTHORS Waterston, R.

COMMENT

TITLE Direct Submission

JOURNAL Submitted (16-APR-2002) Department of Genetics, Washington

University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

On Mar 29, 2002 this sequence version replaced gi:18767626.

----- Genome Center

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: http://genome.wustl.edu/gsc

Contact: sapiens@watson.wustl.edu

----- Summary Statistics Center project name: H NH1413K20

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see http://genome.wustl.edu/gsc

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P.Y., Zhao, B., Frengen, E., Tateno, M., Catanese, J.J. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (http://www.resgen.com) or Pieter de Jong and coworkers at http://www.chori.org

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-489K22, 2000 bp overlap. Actual end is at base position 139342 of RP11-1413K20.

The region between 132012 and 132017 is covered only by a pcr product of clone DNA.

FEATURES

Location/Qualifiers

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misc feature 93. .279

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 Matches
           93; Conservative
                                0; Mismatches
                                                 9; Indels
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Db
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Qy
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Dh
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AC145533
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LOCUS
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DEFINITION
           pieces.
ACCESSION
           AC145533
           AC145533.1 GI:32996774
VERSION
           HTG; HTGS PHASE1; HTGS DRAFT.
KEYWORDS
SOURCE
           Lemur catta (ring-tailed lemur)
  ORGANISM Lemur catta
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Strepsirhini; Lemuridae; Lemur.
REFERENCE
              (bases 1 to 159346)
           Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
 AUTHORS
           Peng, Z., Malinov, I. and Rubin, E.M.
  TITLE
           Direct Submission
  JOURNAL
           Unpublished
           2 (bases 1 to 159346)
REFERENCE
 AUTHORS
           Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
           Peng, Z., Malinov, I. and Rubin, E.M.
           Direct Submission
  TITLE
  JOURNAL
           Submitted (19-JUL-2003) Genome Sciences, Lawrence Berkeley National
           Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
COMMENT
           Draft Sequence Produced by Berkeley PGA
           Web site: http://pga.lbl.gov
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           Center Project Name: L105-138H20
           Bac Clone Name: LB2-138H20
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available at:
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           Funding agent: Programs for Genomic Applications (NHLBI)
           if library name is LB1 to LB4, please see website
           for a description: http://www-gsd.lbl.gov/cheng/BAC.html
           These libraries are available through the BACPAC Resources Center:
           http://www.chori.org/bacpac/libraryres.htm as LBNL-1 to LBNL-4.
           Summary Statistics:
           Sequencing vector: Plasmid; pUC18
           Chemistry: Dye-terminator Big Dye
           Assembly program: Phrap version 0.990329.
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DEFINITION Homo sapiens sterolin-2 (ABCG8) mRNA, complete cds.
ACCESSION
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VERSION
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Additional information on comparative analysis and ordering are

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            1 (bases 1 to 2679)
REFERENCE
  AUTHORS
            Lu, K., Lee, M.H., Hazard, S., Brooks-Wilson, A., Hidaka, H., Kojima, H.,
            Ose, L., Stalenhoef, A.F., Mietinnen, T., Bjorkhem, I., Bruckert, E.,
            Pandya, A., Brewer, H.B. Jr., Salen, G., Dean, M., Srivastava, A. and
            Patel, S.B.
  TITLE
            Two genes that map to the STSL locus cause sitosterolemia: genomic
            structure and spectrum of mutations involving sterolin-1 and
            sterolin-2, encoded by ABCG5 and ABCG8, respectively
            Am. J. Hum. Genet. 69 (2), 278-290 (2001)
  JOURNAL
            21344600
 MEDLINE
   PUBMED
            11452359
            2 (bases 1 to 2679)
REFERENCE
 AUTHORS
            Lu, K., Lee, M.-H. and Patel, S.B.
            Direct Submission
  TITLE
  JOURNAL
            Submitted (29-NOV-2000) Division of Endocrinology, Diabetes and
            Medical Genetics, Medical University of South Carolina, 114 Doughty
            Street, STB541, Charleston, SC 29403, USA
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Qу

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ACCESSION
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           AF351813.1 GI:15146432
VERSION
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           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              (bases 1 to 4665)
REFERENCE
 AUTHORS
           Lu, K., Lee, M.H., Hazard, S., Brooks-Wilson, A., Hidaka, H., Kojima, H.,
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           Pandya, A., Brewer, H.B. Jr., Salen, G., Dean, M., Srivastava, A. and
           Patel, S.B.
  TITLE
           Two genes that map to the STSL locus cause sitosterolemia: genomic
           structure and spectrum of mutations involving sterolin-1 and
           sterolin-2, encoded by ABCG5 and ABCG8, respectively
           Am. J. Hum. Genet. 69 (2), 278-290 (2001)
  JOURNAL
 MEDLINE
           21344600
   PUBMED
           11452359
REFERENCE
           2 (bases 1 to 4665)
 AUTHORS
           Lu, K.
  TITLE
           Direct Submission
           Submitted (21-FEB-2001) Division of Endocrinology, Diabetes and
  JOURNAL
           Medical Genetics, Medical University of South Carolina, 114 Doughty
           St, STB 541, Charleston, SC 29403, USA
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RESULT 17 AC084265/c

LOCUS AC084265 127066 bp DNA linear PRI 11-DEC-2001

DEFINITION Homo sapiens chromosome 2, clone CTB-2367F13, complete sequence.

ACCESSION AC084265

VERSION AC084265.4 GI:17488659

KEYWORDS HTG.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 127066)

AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E. TITLE Homo sapiens chromosome 2, clone CTB-2367F13

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 127066)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., **AUTHORS** Anderson, S., Barna, N., Bastien, V., Beda, F., Boguslavkiy, L., Boukhgalter, B., Brown, A., Burkett, G., Campopiano, A., Castle, A., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Heaford, A., Horton, L., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A., LaRocque, K., Lamazares, R., Landers, T., Lehoczky, J., Levine, R., Lieu, C., Liu, G., Macdonald, P., Marquis, N., McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Morrow, J., Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Olivar, T.M., Oliver, J., Peterson, K., Pierre, N., Pisani, C., Pollara, V., Raymond, C., Rieback, M., Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P., Sougnez, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Tirrell, A., Travers, M., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zimmer, A. and Zody, M.

TITLE Direct Submission

JOURNAL Submitted (18-OCT-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

REFERENCE 3 (bases 1 to 127066)

AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Boguslavkiy, L., Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R., Landers, T., Lehoczky, J., Levine, R., Liu, G.,

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Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (24-AUG-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
   (bases 1 to 127066)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
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Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
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Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (11-DEC-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Dec 11, 2001 this sequence version replaced gi:15284200.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
    Center: Whitehead Institute/ MIT Center for Genome Research
    Center code: WIBR
    Web site: http://www-seq.wi.mit.edu
    Contact: sequence submissions@genome.wi.mit.edu
      ----- Project Information
    Center project name: L11578
    Center clone name: 2367 F 13
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FEATURES source

TITLE

COMMENT

JOURNAL

TITLE

REFERENCE

JOURNAL

AUTHORS

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DEFINITION
           ordered pieces.
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ACCESSION AC146787

VERSION AC146787.1 GI:37497135

KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT.

SOURCE Aotus nancymaae (Ma's night monkey)

ORGANISM Aotus nancymaae

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.

REFERENCE 1 (bases 1 to 178016)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 178016)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Submitted (03-OCT-2003) Genome Sciences, Lawrence Berkeley National

Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA

COMMENT

Sequence Produced by Berkeley PGA

Web site: http://pga.lbl.gov

Center Code: PGABERK
Center Project Name: W010
Bac Clone Name: CH258-323A5

This sequence has been compared to sequences of other species using Vista (http://www-gsd.lbl.gov/VISTA). The results can be viewed at:

http://pga.lbl.gov/cgi-bin/search cvcgd?type=n&value=ABCG5

The order-orientation of the draft sequence was accomplished by using:

Avid (http://baboon.math.berkeley.edu/mavid),

Lagan (http://lagan.stanford.edu/) and paired end information.

Funding agent: Programs for Genomic Applications (NHLBI)

Summary Statistics:

Sequencing vector: Plasmid; pUC18 Chemistry: Dye-terminator Big Dye

Assembly program: Phrap version 0.990329.

- * NOTE: This is a 'working draft' sequence. It currently
- * consists of 4 contigs. Gaps between the contigs
- * are represented as runs of N. The order of the pieces
- * is believed to be correct as given, however the sizes
- * of the gaps between them are based on estimates that have
- * provided by the submittor.
- * This sequence will be replaced
- * by the finished sequence as soon as it is available and
- * the accession number will be preserved.
- * 1 32150: contig of 32150 bp in length
- * 32151 32250: gap of unknown length
- 32251 56222: contig of 23972 bp in length
- * 56223 56322: gap of unknown length
- * 56323 173105: contig of 116783 bp in length
- * 173106 173205: gap of unknown length
- * 173206 178016: contig of 4811 bp in length.

FEATURES

Location/Oualifiers

source

1. .178016

/organism="Aotus nancymaae"
/mol_type="genomic DNA"
/db_xref="taxon:37293"
/clone="CH258-323A5"

ORIGIN

Query Match 82.7%; Score 84.4; DB 2; Length 178016;

Best Local Similarity 89.2%; Pred. No. 1.1e-17;

Matches 91; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

Qy 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60

Db 95879 CTGATAGTTGAGGTCTTTGACCTCCAGGGTATTGGGCTGGCCACTGTAGGTGAAGTACAG 95820

Qy 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102

Db 95819 GCTGTTGTCACTTTCCGAGGAGAACAATCTATCCTGGAGGCC 95778

RESULT 19

AC146466/c

LOCUS AC146466 185045 bp DNA linear HTG 15-AUG-2003

DEFINITION Callithrix jacchus clone CH259-274K20, WORKING DRAFT SEQUENCE, 3

ordered pieces.

ACCESSION AC146466

VERSION AC146466.1 GI:33667132

KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT.

SOURCE Callithrix jacchus (white-tufted-ear marmoset)

ORGANISM Callithrix jacchus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Platyrrhini; Callitrichidae;

Callithrix.

REFERENCE 1 (bases 1 to 185045)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 185045)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Submitted (15-AUG-2003) Genome Sciences, Lawrence Berkeley National

Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA

COMMENT

Sequence Produced by Berkeley PGA

Web site: http://pga.lbl.gov

Center Code: PGABERK Center Project Name: J027 Bac Clone Name: CH259-274K20

This sequence has been compared to sequences of other species using Vista (http://www-gsd.lbl.gov/VISTA). The results can be

viewed at:

http://pqa.lbl.gov/cqi-bin/search cvcqd?type=n&value=ABCG5

```
using:
           Avid (http://baboon.math.berkelev.edu/mavid),
           Lagan (http://lagan.stanford.edu/) and paired end information.
           Funding agent: Programs for Genomic Applications (NHLBI)
           Summary Statistics:
           Sequencing vector: Plasmid; pUC18
           Chemistry: Dye-terminator Big Dye
           Assembly program: Phrap version 0.990329.
           * NOTE: This is a 'working draft' sequence. It currently
           * consists of 3 contigs. Gaps between the contigs
           * are represented as runs of N. The order of the pieces
           * is believed to be correct as given, however the sizes
           * of the gaps between them are based on estimates that have
            * provided by the submittor.
           * This sequence will be replaced
           * by the finished sequence as soon as it is available and
             the accession number will be preserved.
                         49109: contig of 49109 bp in length
                49110
                         49209: gap of unknown length
                49210
                         57420: contig of 8211 bp in length
                57421
                         57520: gap of unknown length
                        185045: contig of 127525 bp in length.
                57521
FEATURES
                    Location/Qualifiers
     source
                    1. .185045
                    /organism="Callithrix jacchus"
                    /mol type="genomic DNA"
                    /db xref="taxon:9483"
                    /clone="CH259-274K20"
ORIGIN
                         82.7%; Score 84.4; DB 2; Length 185045;
 Query Match
                        89.2%; Pred. No. 1.1e-17;
  Best Local Similarity
           91; Conservative
                               0; Mismatches
                                                11;
                                                   Indels
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
             Db
       121318 CTGATAGTTGAGGTCTCTGACCTCCAGGGTATTGGGCTGGCCACTGTAGGTGAAGTACAG
121259
Qу
          61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102
              121258 GCTGTTGTCACTTTCAGAGGAGAACAATCTATCCTGGAGGCC 121217
RESULT 20
AC146464
LOCUS
           AC146464
                                202533 bp
                                             DNA
                                                             HTG 19-AUG-2003
                                                     linear
           Saimiri sciureus clone CH254-84A11, WORKING DRAFT SEQUENCE.
DEFINITION
ACCESSION
           AC146464
VERSION
           AC146464.1 GI:33636782
KEYWORDS
           HTG; HTGS PHASE2; HTGS DRAFT.
SOURCE
           Saimiri sciureus (common squirrel monkey)
 ORGANISM
           Saimiri sciureus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
```

The order-orientation of the draft sequence was accomplished by

```
Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Cebinae;
            Saimiri.
REFERENCE
               (bases 1 to 202533)
            Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
  AUTHORS
            Peng, Z., Malinov, I. and Rubin, E.M.
            Direct Submission
  TITLE
  JOURNAL
            Unpublished
REFERENCE
            2 (bases 1 to 202533)
  AUTHORS
            Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
            Peng, Z., Malinov, I. and Rubin, E.M.
            Direct Submission
  TITLE
            Submitted (14-AUG-2003) Genome Sciences, Lawrence Berkeley National
  JOURNAL
            Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
               (bases 1 to 202533)
REFERENCE
            Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
 AUTHORS
            Peng, Z., Malinov, I. and Rubin, E.M.
 TITLE
            Direct Submission
  JOURNAL
            Submitted (19-AUG-2003) Genome Sciences, Lawrence Berkeley National
            Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
COMMENT
            Sequence Produced by Berkeley PGA
            Web site: http://pga.lbl.gov
            Center Code: PGABERK
            Center Project Name: S030
            Bac Clone Name: CH254-84A11
            This sequence has been compared to sequences of other species
            using Vista (http://www-gsd.lbl.gov/VISTA). The results can be
            viewed at:
            http://pqa.lbl.qov/cqi-bin/search cvcqd?type=n&value=ABCG5
            The order-orientation of the draft sequence was accomplished by
            using:
            Avid (http://baboon.math.berkeley.edu/mavid),
            Lagan (http://lagan.stanford.edu/) and paired end information.
            Funding agent: Programs for Genomic Applications (NHLBI)
            Summary Statistics:
            Sequencing vector: Plasmid; pUC18
            Chemistry: Dye-terminator Big Dye
            Assembly program: Phrap version 0.990329.
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 1 contigs. Gaps between the contigs
            * are represented as runs of N. The order of the pieces
            * is believed to be correct as given, however the sizes
            * of the gaps between them are based on estimates that have
            * provided by the submittor.
            * This sequence will be replaced
            * by the finished sequence as soon as it is available and
            * the accession number will be preserved.
                         202533: contig of 202533 bp in length.
                     1
FEATURES
                     Location/Qualifiers
                     1. .202533
     source
                     /organism="Saimiri sciureus"
                     /mol type="genomic DNA"
                     /db xref="taxon:9521"
```

ORIGIN

```
81.2%; Score 82.8; DB 2; Length 202533;
  Query Match
                         88.2%; Pred. No. 4e-17;
  Best Local Similarity
                                                               0; Gaps
                                                                           0;
           90; Conservative
                               0; Mismatches
                                               12;
                                                     Indels
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
             27346 CTGATAGTTGAGGTCTTTGACCTCCAGGGTATTGGGCTGGCCACTGTAGGTGAAGTACAG 27405
Db
          61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102
QУ
              27406 GCTGTTGTCACTTTCGGAGGAGAACAATCTATCCTGGAGGCC 27447
Db.
RESULT 21
AC146286/c
                                 207760 bp
                                             DNA
                                                     linear
                                                              HTG 15-AUG-2003
LOCUS
           AC146286
           Callicebus moloch clone LB5-414K16, WORKING DRAFT SEQUENCE, 2
DEFINITION
           ordered pieces.
ACCESSION
           AC146286
VERSION
           AC146286.2 GI:33667134
           HTG; HTGS PHASE2; HTGS DRAFT.
KEYWORDS
           Callicebus moloch (Dusky titi)
SOURCE
           Callicebus moloch
  ORGANISM
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Callicebinae;
           Callicebus.
REFERENCE
           1 (bases 1 to 207760)
  AUTHORS
           Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
           Peng, Z., Malinov, I. and Rubin, E.M.
  TITLE
           Direct Submission
  JOURNAL
           Unpublished
REFERENCE
           2
              (bases 1 to 207760)
           Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
  AUTHORS
           Peng, Z., Malinov, I. and Rubin, E.M.
  TITLE
           Direct Submission
  JOURNAL
           Submitted (02-AUG-2003) Genome Sciences, Lawrence Berkeley National
           Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
REFERENCE
              (bases 1 to 207760)
           Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
  AUTHORS
           Peng, Z., Malinov, I. and Rubin, E.M.
  TITLE
           Direct Submission
           Submitted (15-AUG-2003) Genome Sciences, Lawrence Berkeley National
  JOURNAL
           Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
           On Aug 15, 2003 this sequence version replaced gi:33413351.
COMMENT
           Sequence Produced by Berkeley PGA
           Web site: http://pga.lbl.gov
           Center Code: PGABERK
           Center Project Name: T039
           Bac Clone Name: LB5-414K16
```

This sequence has been compared to sequences of other species using Vista (http://www-gsd.lbl.gov/VISTA). The results can be viewed at:

The order-orientation of the draft sequence was accomplished by using: Avid (http://baboon.math.berkeley.edu/mavid), Lagan (http://lagan.stanford.edu/) and paired end information. Funding agent: Programs for Genomic Applications (NHLBI) If the Bac Library Name is LB1 to LB4, please see website for the description: http://www-gsd.lbl.gov/cheng/BAC.html These libraries are available through the BACPAC Resources Center: http://www.chori.org/bacpac/libraryres.htm as LBNL-1 to LBNL-4. Summary Statistics: Sequencing vector: Plasmid; pUC18 Chemistry: Dye-terminator Big Dye Assembly program: Phrap version 0.990329. * NOTE: This is a 'working draft' sequence. It currently * consists of 2 contigs. Gaps between the contigs * are represented as runs of N. The order of the pieces * is believed to be correct as given, however the sizes * of the gaps between them are based on estimates that have * provided by the submittor. * This sequence will be replaced * by the finished sequence as soon as it is available and * the accession number will be preserved. 74764: contig of 74764 bp in length 74864: gap of unknown length 74765 74865 207760: contig of 132896 bp in length. Location/Qualifiers FEATURES 1. .207760 source /organism="Callicebus moloch" /mol type="genomic DNA" /db xref="taxon:9523" /clone="LB5-414K16" ORIGIN 81.2%; Score 82.8; DB 2; Length 207760; Query Match 88.2%; Pred. No. 4e-17; Best Local Similarity 90; Conservative 0; Mismatches 12; Indels Gaps 0; 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Qу 145281 CTGATAGTTGAGGTCTCTGACCTCTAGGGTATTGGGCTGGCCACTGTAGGTGAAGTACAG Db 145222 61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102 Qy 145221 GCTGTTGTCACTTTCAGAGGAGAACAATCTATCCTGGAGGCC 145180 Db RESULT 22 AC146282/c HTG 02-AUG-2003 135280 bp DNA linear LOCUS Takifugu rubripes clone MRC-186C24, WORKING DRAFT SEQUENCE, 7 DEFINITION unordered pieces.

http://pga.lbl.gov/cgi-bin/search cvcgd?type=n&value=ABCG5

```
ACCESSION
            AC146282
VERSION
            AC146282.1 GI:33413347
KEYWORDS
            HTG; HTGS PHASE1; HTGS DRAFT.
SOURCE
            Takifugu rubripes (Fugu rubripes)
 ORGANISM
           Takifugu rubripes
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
            Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
            Tetradontoidea; Tetraodontidae; Takifugu.
            1 (bases 1 to 135280)
REFERENCE
 AUTHORS
            Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
            Peng, Z., Malinov, I. and Rubin, E.M.
            Direct Submission
 TITLE
  JOURNAL
            Unpublished
REFERENCE
            2 (bases 1 to 135280)
 AUTHORS
            Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,
            Peng, Z., Malinov, I. and Rubin, E.M.
  TITLE
            Direct Submission
            Submitted (02-AUG-2003) Genome Sciences, Lawrence Berkeley National
  JOURNAL
            Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
            Draft Sequence Produced by Berkeley PGA
COMMENT
            Web site: http://pga.lbl.gov
            Center Code: PGABERK
            Center Project Name: F069-186C24
            Bac Clone Name: MRC-186C24
            Additional information on comparative analysis and ordering are
            available at:
            http://pga.lbl.gov/cgi-bin/search cvcgd?type=n&value=
            Funding agent: Programs for Genomic Applications (NHLBI)
            Summary Statistics:
            Sequencing vector: Plasmid; pUC18
            Chemistry: Dye-terminator Big Dye
            Assembly program: Phrap version 0.990329.
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 7 contigs. The true order of the pieces
            * is not known and their order in this sequence record is
            * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
            * be preserved.
                     1
                          28849: contig of 28849 bp in length
                 28850
                          28949: gap of unknown length
                          40654: contig of 11705 bp in length
                 28950
                 40655
                          40754: gap of unknown length
                 40755
                          55789: contig of 15035 bp in length
                 55790
                          55889: gap of unknown length
                          70983: contig of 15094 bp in length
                 55890
                          71083: gap of unknown length
                 70984
                          90702: contig of 19619 bp in length
                 71084
                 90703
                          90802: gap of unknown length
                 90803
                         112817: contig of 22015 bp in length
                         112917: gap of unknown length
                112818
                         135280: contig of 22363 bp in length.
                112918
                     Location/Qualifiers
FEATURES
                     1. .135280
     source
```

```
/organism="Takifugu rubripes"
/mol_type="genomic DNA"
/db_xref="taxon:31033"
/clone="MRC-186C24"
```

ORIGIN

Query Match 51.4%; Score 52.4; DB 2; Length 135280; Best Local Similarity 75.6%; Pred. No. 8.3e-07; 0; Mismatches 65; Conservative 21; Indels 0; Gaps 0: Qу 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60 Db 34807 CTCATAGTTGAGGTCGTTGACCTCCAGCTGGTTGCACCCTCCACTGTAGGTGAAGTAGAG 34748 61 ACTGTTGTCACTTTCCGAGGAGAACA 86 Qу 34747 GCTGCTGTCTTCTTCAGTGGAGAACA 34722 Db

RESULT 23 AL928999

LOCUS AL928999 169570 bp DNA linear VRT 24-DEC-2002 DEFINITION Zebrafish DNA sequence from clone CH211-227C6, complete sequence.

ACCESSION AL928999

VERSION AL928999.4 GI:26788223

KEYWORDS HTG.

SOURCE Danio rerio (zebrafish)

ORGANISM Danio rerio

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi;

Cypriniformes; Cyprinidae; Danio.

REFERENCE 1 (bases 1 to 169570)

AUTHORS Heath, P.

TITLE Direct Submission

JOURNAL Submitted (21-DEC-2002) Wellcome Trust Sanger Institute, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquiries: zface@sanger.ac.uk

Clone requests: clonerequest@sanger.ac.uk

COMMENT On Dec 13, 2002 this sequence version replaced gi:25055310.

----- Genome Center

Center: Wellcome Trust Sanger Institute

Center code: SC

Web site: http://www.sanger.ac.uk

Contact: zface@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare

```
occasion of the clone being a YAC.
           The following abbreviations are used to associate primary accession
           numbers given in the feature table with their source databases:
           Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information
           on the WORMPEP database can be found at
           http://www.sanger.ac.uk/Projects/C elegans/wormpep Repeat names
           beginning 'Dr' were identified by the Recon repeat discovery system
           (Zhirong Bao and Sean Eddy, submitted), and those beginning 'drr'
           were identified by Rick Waterman (Stephen Johnson lab, WashU). For
           further information see http://www/Projects/D rerio/fishmask.shtml
           CH211-227C6 is from a CHORI-211 BAC library
           VECTOR: pTARBAC2.1.
                   Location/Qualifiers
FEATURES
                   1. .169570
    source
                   /organism="Danio rerio"
                   /mol type="genomic DNA"
                   /db xref="taxon:7955"
                   /clone="CH211-227C6"
                   /clone lib="CHORI-211"
ORIGIN
                        35.1%; Score 35.8; DB 5; Length 169570;
 Query Match
 Best Local Similarity
                        63.2%; Pred. No. 0.36;
                                                                 Gaps
                                                                        0;
           55; Conservative
                               0; Mismatches
                                               32;
                                                   Indels
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
                +11
                                                   Db
          61 ACTGTTGTCACTTTCCGAGGAGAACAA 87
Qν
             74407 ACTGCTGTCCTCTCTGGAGATGAAAA 74433
Db
RESULT 24
BX004832/c
                                                    linear
                                                            VRT 25-NOV-2003
           BX004832
                                190952 bp
                                            DNA
LOCUS
           Zebrafish DNA sequence from clone CH211-89M19, complete sequence.
DEFINITION
ACCESSION
           BX004832
VERSION
           BX004832.9 GI:38524388
KEYWORDS
           HTG.
           Danio rerio (zebrafish)
SOURCE
  ORGANISM
           Danio rerio
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
           Cypriniformes; Cyprinidae; Danio.
           1 (bases 1 to 190952)
REFERENCE
           Harrison, E.
  AUTHORS
           Direct Submission
  TITLE
           Submitted (25-NOV-2003) Wellcome Trust Sanger Institute, Hinxton,
  JOURNAL
           Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
           zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
           On Nov 25, 2003 this sequence version replaced gi:31335509.
COMMENT
           ---- Genome Center
           Center: Wellcome Trust Sanger Institute
           Center code: SC
           Web site: http://www.sanger.ac.uk
```

Contact: zfish-help@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep Repeat names beginning 'Dr' were identified by the Recon repeat discovery system (Zhirong Bao and Sean Eddy, submitted), and those beginning 'drr' were identified by Rick Waterman (Stephen Johnson lab, WashU). For further information see

http://www.sanger.ac.uk/Projects/D_rerio/fishmask.shtml CH211-89M19 is from a CHORI-211 BAC library

VECTOR: pTARBAC2.1

Clone-derived Zebrafish pUC subclones occasionally display inconsistency over the length of mononucleotide A/T runs and conserved TA repeats. Where this is found the longest good quality representation will be submitted.

FEATURES

Location/Qualifiers

source

1. .190952

/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="CH211-89M19"
/clone lib="CHORI-211"

ORIGIN

Query Match 33.5%; Score 34.2; DB 5; Length 190952;

Best Local Similarity 62.1%; Pred. No. 1.3;

Matches 54; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

Qy 1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60

Qy 61 ACTGTTGTCACTTTCCGAGGAGAACAA 87

Db 92028 ACTGCTGTCCTCTGGAGATGAAAA 92002

RESULT 25 BX571838

LOCUS BX571838 226929 bp DNA linear HTG 27-SEP-2003

```
Danio rerio clone DKEY-205N7, WORKING DRAFT SEQUENCE, 14 unordered
DEFINITION
           pieces.
ACCESSION
           BX571838
VERSION
           BX571838.3 GI:36796624
KEYWORDS
           HTG; HTGS PHASE1; HTGS DRAFT; HTGS FULLTOP.
           Danio rerio (zebrafish)
SOURCE
 ORGANISM Danio rerio
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
           Cypriniformes; Cyprinidae; Danio.
REFERENCE
           1 (bases 1 to 226929)
 AUTHORS
           Mclaren, S.
           Direct Submission
 TITLE
 JOURNAL
           Submitted (26-SEP-2003) Wellcome Trust Sanger Institute, Hinxton,
           Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
           zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
           On Sep 27, 2003 this sequence version replaced gi:33386624.
COMMENT
           ----- Genome Center
           Center: Wellcome Trust Sanger Institute
           Center code: SC
           Web site: http://www.sanger.ac.uk
           Contact: zfish-help@sanger.ac.uk
            ----- Project Information
           Center project name: zK205N7
            ----- Summary Statistics
           Assembly program: XGAP4; version 4.5
           Chemistry: Dye-terminator; 100% of reads
           Consensus quality: 223662 bases at least Q40
           Consensus quality: 224399 bases at least Q30
           Consensus quality: 224947 bases at least Q20
           Insert size: 225629; sum-of-contigs
           Insert size: 196940; 4.8% error; agarose-fp
           Quality coverage: 6.66x in Q20 bases; sum-of-contigs Quality
           coverage: 7.66x in Q20 bases; agarose-fp
           -----
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 14 contigs. The true order of the pieces
            * is not known and their order in this sequence record is
            * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
            * be preserved.
                    1
                         10067: contig of 10067 bp in length
                 10068
                         10167: gap of 100 bp
                 10168
                         24021: contig of 13854 bp in length
                24022
                         24121: gap of 100 bp
                         28447: contig of 4326 bp in length
                24122
                         28547: gap of 100 bp
                28448
                         47699: contig of 19152 bp in length
                28548
                         47799: gap of 100 bp
                 47700
                         68972: contig of 21173 bp in length
                 47800
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                 68973
                 69073
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                         74019: gap of 100 bp
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106234: contig of 32215 bp in length

74020

106235 106334: gap of 100 bp

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106335
                         126675: contig of 20341 bp in length
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                         126775: gap of 100 bp
                126776
                         145072: contig of 18297 bp in length
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                145173
                         161897: contig of 16725 bp in length
                161898
                         161997: gap of 100 bp
                161998
                         198188: contig of 36191 bp in length
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                         198288: gap of 100 bp
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                         204891: contig of 6603 bp in length
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                         204991: gap of 100 bp
                204992
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Db
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ACCESSION
           M55034
VERSION
           M55034.1 GI:180298
KEYWORDS
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SOURCE
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REFERENCE
              (bases 1 to 203)
  AUTHORS
           Kerem, B.-S., Zielenski, J., Markiewicz, D., Bozon, D., Gazit, E.,
           Yahav, J., Kennedy, D., Riordan, J.R., Collins, F.S., Rommens, J.M. and
           Tsui, L.-C.
           Identification of mutations in regions corresponding to the two
  TITLE
           putative nucleotide (ATP)-binding folds of the cystic fibrosis gene
  JOURNAL
           Proc. Natl. Acad. Sci. U.S.A. 87 (21), 8447-8451 (1990)
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DEFINITION
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            (CFTR) gene, exon 10.
ACCESSION
           M55025
VERSION
           M55025.1 GI:180297
KEYWORDS
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           Homo sapiens (human)
SOURCE
  ORGANISM
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REFERENCE
           1 (bases 1 to 206)
 AUTHORS
           Kerem, B.-S., Zielenski, J., Markiewicz, D., Bozon, D., Gazit, E.,
           Yahav, J., Kennedy, D., Riordan, J.R., Collins, F.S., Rommens, J.M. and
           Tsui, L.-C.
  TITLE
           Identification of mutations in regions corresponding to the two
           putative nucleotide (ATP)-binding folds of the cystic fibrosis gene
  JOURNAL
           Proc. Natl. Acad. Sci. U.S.A. 87 (21), 8447-8451 (1990)
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Db
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Qу
             88 TTCTCAGTTTTCCTGGA 104
Db
RESULT 28
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LOCUS
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DEFINITION
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           AF162161
ACCESSION
           AF162161.1 GI:8886448
VERSION
KEYWORDS
SEGMENT
           11 of 27
           Macaca fascicularis (crab-eating macague)
SOURCE
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           Mammalia; Eutheria; Primates; Catarrhini; Cercopithecidae;
           Cercopithecinae; Macaca.
           1 (bases 1 to 261)
REFERENCE
 AUTHORS
           Wine, J.J., Kuo, E., Hurlock, G., Glavac, D. and Dean, M.
 TITLE
           Genomic sequence of CFTR in five primate species
 JOURNAL
           Unpublished
REFERENCE
           2 (bases 1 to 261)
 AUTHORS
           Wine, J.J., Kuo, E., Hurlock, G., Glavac, D. and Dean, M.
 TITLE
           Direct Submission
 JOURNAL
           Submitted (24-JUN-1999) Psychology, Stanford University, Building
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Db
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Qy
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DEFINITION
            (CFTR) gene, exon 10.
            AF162357
ACCESSION
            AF162357.1 GI:5679203
VERSION
KEYWORDS
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SEGMENT
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SOURCE
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REFERENCE
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  AUTHORS
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  TITLE
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REFERENCE
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Db
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Qу
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          111 TTCTCAGTTTTCCTGGA 127
Db
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MNCFTR11
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                                               DNA
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LOCUS
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            Macaca nemestrina cystic fibrosis transmembrane conductance
DEFINITION
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ACCESSION
            AF162384
            AF162384.1 GI:5679232
VERSION
KEYWORDS
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11 of 27
SEGMENT
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SOURCE
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           Mammalia; Eutheria; Primates; Catarrhini; Cercopithecidae;
           Cercopithecinae; Macaca.
           1 (bases 1 to 261)
REFERENCE
           Wine, J.J., Kuo, E., Hurlock, G., Glavac, D. and Dean, M.
 AUTHORS
           CFTR genomic sequences from five primate species
 TITLE
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  JOURNAL
           2 (bases 1 to 261)
REFERENCE
           Wine, J.J., Kuo, E., Hurlock, G., Glavac, D. and Dean, M.
 AUTHORS
           Direct Submission
 TITLE
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  JOURNAL
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DEFINITION
            regulator (CFTR) gene, exon 10.
ACCESSION
           AF162411
           AF162411.1 GI:5679263
VERSION
KEYWORDS
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SEGMENT
SOURCE
            Papio anubis (olive baboon)
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           Mammalia; Eutheria; Primates; Catarrhini; Cercopithecidae;
            Cercopithecinae; Papio.
REFERENCE
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           Wine, J.J., Kuo, E., Hurlock, G., Glavac, D. and Dean, M.
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  TITLE
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  JOURNAL
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REFERENCE
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Qy
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Db
RESULT 32
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                                                                PRI 18-APR-1998
LOCUS
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            Macaca mulatta cystic fibrosis transmembrane conductance regulator
DEFINITION
            (CFTR) gene, exon 10.
ACCESSION
            AF016934
VERSION
            AF016934.1 GI:3057098
KEYWORDS
SEGMENT
            11 of 27
SOURCE
            Macaca mulatta (rhesus monkey)
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            1 (bases 1 to 261)
REFERENCE
            Wine, J.J., Glavac, D., Hurlock, G., Robinson, C., Lee, M., Potocnik, U.,
  AUTHORS
            Ravnik-Glavac, M. and Dean, M.
            Genomic DNA sequence of Rhesus (M. mulatta) cystic fibrosis (CFTR)
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            Mamm. Genome 9 (4), 301-305 (1998)
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RESULT 33
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DEFINITION
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VERSION
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KEYWORDS
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SOURCE
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           1 (bases 1 to 420)
REFERENCE
           Mitchell, L.G. and Garcia-Blanco, M.A.
  AUTHORS
           Methods and compositions for use in spliceosome mediated RNA
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           Patent: US 6280978-A 64 28-AUG-2001;
  JOURNAL
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ACCESSION
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REFERENCE
            Gordon, J. and Rundell, C.A.
  AUTHORS
            Compositions and methods relating to control DNA construct
  TITLE
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LOCUS
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DEFINITION
            gene, exon 10.
            M55115
ACCESSION
            M55115.1 GI:306520
VERSION
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KEYWORDS
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            1 (bases 1 to 831)
REFERENCE
            Zielenski, J., Rozmahel, R., Bozon, D., Kerem, B., Grzelczak, Z.,
  AUTHORS
            Riordan, J.R., Rommens, J. and Tsui, L.C.
            Genomic DNA sequence of the cystic fibrosis transmembrane
  TITLE
            conductance regulator (CFTR) gene
            Genomics 10 (1), 214-228 (1991)
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JOURNAL

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Db
RESULT 36
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DEFINITION
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ACCESSION
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VERSION
KEYWORDS
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           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
              (bases 1 to 831)
 AUTHORS
           Bouffard, G.G., Iyer, L.M., Idol, J.R., Braden, V.V., Cunningham, A.F.,
           Weintraub, L.A., Mohr-Tidwell, R.M., Peluso, D.C., Fulton, R.S.,
           Leckie, M.P. and Green, E.D.
           A collection of 1814 human chromosome 7-specific STSs
 TITLE
           Genome Res. 7 (1), 59-64 (1997)
  JOURNAL
           97189344
  MEDLINE
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           9037602
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REFERENCE
           2
  AUTHORS
           Green, E.D.
           Human chromosome 7 STSs (1997)
  TITLE
  JOURNAL
           Unpublished (1997)
COMMENT
           Synonyms: CFTR
           GDB: GDB: 3754054
           GDB DSEG: CFTR
           Contact: Eric D. Green
           Genome Technology Branch
           National Human Genome Research Institute/NIH
```

91257831

```
Tel: 3014020201
           Fax: 3014024735
           Email: egreen@nhgri.nih.gov
           Primer A: CAGTTTTCCTGGATTATGCCTGG
           Primer B: GTTGGCATGCTTTGATGACGCTTC
           STS size: 100
           PCR Profile:
                   Presoak:
                                    0 degrees C for 0.00 minute(s)
                   Denaturation:
                                   92 degrees C for
                                                     1.00 \text{ minute(s)}
                                   62 degrees C for
                                                     2.00 minute(s)
                   Annealing:
                   Polymerization: 72 degrees C for 2.00 minute(s)
                   PCR Cycles:
                                   35
                   Thermal Cycler: PerkinElmer TC
           Protocol:
                                    30-100 ng
                   Template:
                   Primer:
                                    each 1 uM
                   dNTPs:
                                   each 200 uM
                   Tag Polymerase: 0.05 units/ul
                   Total Vol:
                                    5 ul
           Buffer:
                   MqCl2:
                                    2.5 mM
                                    50 mM
                   KCl:
                                    10 mM
                   Tris-HCl:
                                    8.3
                   :Hq
             This STS was developed from sequence determined by
                           See GenBank record: M55115
                                                        For additional
           investigator.
           information about the NHGRI chromosome 7 mapping project, see
           http://www.nhqri.nih.qov/DIR/GTB/CHR7. Also see Genomics
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Db
Qу
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49 Convent Dr., MSC4431, Bldg. 49, Rm. 2A08, Bethesda, MD 20892

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           Welsh, M.J. and Sheppard, D.N.
  AUTHORS
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REFERENCE
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           Welsh, M.J. and Sheppard, D.N.
  AUTHORS
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ACCESSION
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REFERENCE
            1 (bases 1 to 2908)
  AUTHORS
           Xu, Z. and Gruenert, D.C.
            Human CFTR gene sequences in regions flanking exon 10: a simple
  TITLE
            repeat sequence polymorphism in intron 9
            Biochem. Biophys. Res. Commun. 219 (1), 140-145 (1996)
  JOURNAL
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            96190683
   PUBMED
            8619797
            Original source text: Homo sapiens (clone: T6/20) DNA.
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REFERENCE
          Teem, J.L.
 AUTHORS
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          Patent: US 6468793-A 1 22-OCT-2002;
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 AUTHORS
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 AUTHORS
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          Patent: US 6468793-A 9 22-OCT-2002;
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REFERENCE
 AUTHORS
           Teem, J.L.
 TITLE
           Materials and method for detecting interaction of cftr polypeptides
 JOURNAL
           Patent: WO 0125421-A 3 12-APR-2001;
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4: geneseqn2001as:* 5: geneseqn2001bs:*

6: geneseqn2002s:*

7: geneseqn2003as:*

8: geneseqn2003bs:*

9: genesegn2003cs:*

10: geneseqn2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.		Score	<pre>% Query Match</pre>	Length	DB	ID Description	
С	1	102	100.0	2019	7	AAD48881	Aad48881 Mouse ABC
С	2	102	100.0	2564	6	ABN90022	Abn90022 Mouse clo
	3	102	100.0	6043	7	AAD48884	Aad48884 ABCG5-ABC
С	4	87.6	85.9	2669	7	AAD48883	Aad48883 Human ABC
	5	32.2	31.6	180	4	ABA71163	Aba71163 Human foe
	6	32.2	31.6	180	4	AAI51393	Aai51393 Probe #20
	7	32.2	31.6	180	4	AAK45448	Aak45448 Human bon

8 .	32.2	31.6	180	4	AAK19459	Aak19459	Human bra
9	32.2	31.6	180	4	ABS45131	Abs45131	Human liv
10	32.2	31.6	180	6	ABS19713	Abs19713	Human gen
11	32.2	31.6	494	4	ABA58823	Aba58823	Human foe
12	32.2	31.6	494	4	AAI38528	Aai38528	Probe #72
13	32.2	31.6	494	4	AAK32713	Aak32713	Human bon
14	32.2	31.6	494	4	AAK06977	Aak06977	Human bra
15	32.2	31.6	494	4	ABS32432	Abs32432	Human liv
16	32.2	31.6	494	6	ABS07509	Abs07509	Human gen
17	32.2	31.6	500	3	AAZ99413	Aaz99413	Trans-spl
18	32.2	31.6	500	6	ABQ73502	Abq73502	Pre-trans
19	32.2	31.6	795	7	ABZ24468	Abz24468	Cystic fi
20	32.2	31.6	831	9	ADE77694	Ade77694	Human cys
21	32.2	31.6	2640	2	AAT04005	Aat04005	Truncated
22	32.2	31.6	3069	6	ABQ73521	Abq73521	Mouse fac
23	32.2	31.6	4443	4	AAF84742	. Aaf84742	DNA encod
24	32.2	31.6	4443	8	ABX16100	Abx16100	Human cDN
25	32.2	31.6	4443	8	ABX16094	Abx16094	Human cDN
26	32.2	31.6	4443	8	ABX16099	Abx16099	Human cDN
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28	32.2	31.6	4443	8	ABX16103	Abx16103	Human cDN
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47	32.2	31.6	6129	2	AAQ13071		CFTR 1717
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49	32.2	31.6	6129	2	AAQ13065		CFTR L107
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ALIGNMENTS

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RESULT 1
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ID AAD48881 standard; DNA; 2019 BP.
XX
AC AAD48881;
XX
DT 24-MAR-2003 (first entry)
XX
DE Mouse ABCG8 DNA.
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XX
    ABC family cholesterol transporter; ABCG8; sterol-related disorder;
ΚW
KW
     sitosterolaemia; hyperlipidaemia; hypercholesterolaemia; gall stone;
     HDL deficiency; atherosclerosis; nutritional deficiency; gene therapy;
KW
     mouse; ATP-binding cassette; sitosterolaemia susceptibility gene; SSG;
KW
    ABCG5; gene; ds.
KW
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    Mus sp.
XX
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XX
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XX
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    20-NOV-2001; 2001WO-US043823.
XX
PR
    20-NOV-2000; 2000US-0252235P.
PR
     28-NOV-2000; 2000US-0253645P.
XX
     (TULA-) TULARIK INC.
PA
PA
     (TEXA ) UNIV TEXAS SYSTEM.
XX
PΙ
    Hobbs HH, Shan B, Barnes R, Tian H;
XX
DR
    WPI; 2003-058548/05.
DR
     P-PSDB; AAE31703.
ХX
PT
    New ABCG8 polypeptides and nucleic acids, useful for treating sterol-
PT
     related disorders e.g. sitosterolemia, hypercholesterolemia,
PT
     hyperlipidemia, gall stones, HDL deficiency, atherosclerosis, or
    nutritional deficiencies.
PT
XX
PS
    Claim 13; Page 75; 94pp; English.
XX
    The invention relates to ATP-binding cassette (ABC) family cholesterol
CC
CC
    transporter, ABCG8 polypeptides and polynucleotides. The invention also
    provides ABCG5 polypeptides and polynucleotides. ABCG5 gene is also known
CC
     as sitosterolaemia susceptibility gene (SSG). Sequences of the invention
CC
    are useful for treating or preventing sterol-related disorders such as
CC
    sitosterolaemia, hyperlipidaemia, hypercholesterolaemia, gall stones, HDL
CC
    deficiency, atherosclerosis and nutritional deficiencies. They are also
CC
CC
     useful in gene therapy. The present sequence is mouse ABCG8 DNA
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                                0; Mismatches
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XX
DT
     16-AUG-2002 (first entry)
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DE
     Mouse clone IMX3 67 extended sequence.
XX
KW
     Mouse; antiinflammatory; gene therapy; ileitis; DST; ss; TOGA;
KW
     digital sequence tag; total gene expression analysis.
XX
OS
     Mus musculus.
XX
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     WO200231114-A2.
XX
PD
    18-APR-2002.
XX
PF
     11-OCT-2001; 2001WO-US032091.
XX
     11-OCT-2000; 2000US-0239483P.
PR
XX
PA
     (DIGI-) DIGITAL GENE TECHNOLOGIES INC.
XX
PΙ
     Viney JL, Sims JE, Dubose RF, Baum PR,
                                               Hasel KW,
                                                          Hilbush BS;
XX
DR
     WPI; 2002-426279/45.
XX
     New isolated nucleic acid molecules that are associated with ileitis, for
PT
PΤ
     preventing, treating, modulating and diagnosing ileitis in a mammalian
PT
     subject.
XX
PS
     Claim 1; Page 266-268; 273pp; English.
XX
CC
     The invention relates to a novel isolated nucleic acid molecule
     comprising a polynucleotide having one of 90 polynucleotide sequences,
CC
CC
     given in the specification. The polynucleotides of the invention have
CC
     antiinflammatory activity, and may have a use in gene therapy. The
CC
     polynucleotide or a polypeptide encoded by it is used for preventing,
CC
     treating, modulating or ameliorating a medical condition such as ileitis.
CC
     The polypeptide or polynucleotide is also useful for manufacturing a
     medicament for treating ileitis. The sequence represents a an extended
CC
CC
     cDNA digital sequence tag obtained from a mouse clone by the TOGA (total
CC
     gene expression analysis) method
XX
SQ
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  Best Local Similarity
                         100.0%; Pred. No. 2.9e-25;
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                               0; Mismatches
                                                 0; Indels
                                                                            0:
                                                                0; Gaps
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Qy
             Db
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    AAD48884;
XX
    24-MAR-2003 (first entry)
DT
XX
DE
    ABCG5-ABCG8 DNA.
XX
KW
    ABC family cholesterol transporter; ABCG8; sterol-related disorder;
    sitosterolaemia; hyperlipidaemia; hypercholesterolaemia; gall stone;
ΚW
    HDL deficiency; atherosclerosis; nutritional deficiency; gene therapy;
KW
    ATP-binding cassette; sitosterolaemia susceptibility gene; SSG; ABCG5;
KW
KW
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XX
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    20-NOV-2001; 2001WO-US043823.
XX
PR
    20-NOV-2000; 2000US-0252235P.
    28-NOV-2000; 2000US-0253645P.
PR
XX
PA
     (TULA-) TULARIK INC.
PΑ
     (TEXA ) UNIV TEXAS SYSTEM.
XX
PI
    Hobbs HH,
               Shan B, Barnes R, Tian H;
XX
    WPI; 2003-058548/05.
DR
XX
    New ABCG8 polypeptides and nucleic acids, useful for treating sterol-
PT
     related disorders e.g. sitosterolemia, hypercholesterolemia,
PT
PT
    hyperlipidemia, gall stones, HDL deficiency, atherosclerosis, or
PT
    nutritional deficiencies.
XX
    Disclosure; Fig 3; 94pp; English.
PS
XX
CC
    The invention relates to ATP-binding cassette (ABC) family cholesterol
CC
    transporter, ABCG8 polypeptides and polynucleotides. The invention also
CC
    provides ABCG5 polypeptides and polynucleotides. ABCG5 gene is also known
    as sitosterolaemia susceptibility gene (SSG). Sequences of the invention
CC
    are useful for treating or preventing sterol-related disorders such as
CC
    sitosterolaemia, hyperlipidaemia, hypercholesterolaemia, gall stones, HDL
CC
    deficiency, atherosclerosis and nutritional deficiencies. They are also
CC
CC
    useful in gene therapy. The present sequence is ABCG8- ABCG5 DNA
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                                                    Indels
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                                                                          0;
                                                                  Gaps
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Qу
             3 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 62
Db
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    AAD48883;
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DT
     24-MAR-2003
                  (first entry)
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DE
    Human ABCG8 DNA.
XX
KW
     ABC family cholesterol transporter; ABCG8; sterol-related disorder;
KW
     sitosterolaemia; hyperlipidaemia; hypercholesterolaemia; gall stone;
KW
     HDL deficiency; atherosclerosis; nutritional deficiency; gene therapy;
     human; ATP-binding cassette; sitosterolaemia susceptibility gene; SSG;
KW
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     20-NOV-2000; 2000US-0252235P.
PR
     28-NOV-2000; 2000US-0253645P.
XX
PA
     (TULA-) TULARIK INC.
     (TEXA ) UNIV TEXAS SYSTEM.
PA
XX
PΙ
     Hobbs HH, Shan B, Barnes R, Tian H;
XX
DR
     WPI; 2003-058548/05.
DR
     P-PSDB; AAE31705.
XX
PT
     New ABCG8 polypeptides and nucleic acids, useful for treating sterol-
PT
     related disorders e.g. sitosterolemia, hypercholesterolemia,
PT
     hyperlipidemia, gall stones, HDL deficiency, atherosclerosis, or
PT
     nutritional deficiencies.
XX
PS
     Claim 13; Page 80; 94pp; English.
XX
     The invention relates to ATP-binding cassette (ABC) family cholesterol
CC
CC
     transporter, ABCG8 polypeptides and polynucleotides. The invention also
CC
     provides ABCG5 polypeptides and polynucleotides. ABCG5 gene is also known
     as sitosterolaemia susceptibility gene (SSG). Sequences of the invention
CC
     are useful for treating or preventing sterol-related disorders such as
CC
CC
     sitosterolaemia, hyperlipidaemia, hypercholesterolaemia, gall stones, HDL
CC
     deficiency, atherosclerosis and nutritional deficiencies. They are also
CC
     useful in gene therapy. The present sequence is human ABCG8 DNA
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QУ
                              111411 1114 1114 11141111141 1114 1114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 114 
                     264 CTGGTAGTTGAGGTCTCTGACCTCCAGGGTGTTGGGCTGGCCACTGTAGGTGAAGTACAG 205
Db
                       61 ACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGCC 102
Qу
                                204 GCTGTTGTCACTTTCAGAGGAGAACAATCTATCCTGGAGGCC 163
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          ABA71163;
XX
          01-FEB-2002 (first entry)
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XX
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DE
XX
          Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
KW
XX
OS
          Homo sapiens.
XX
          WO200157277-A2.
PN
XX
          09-AUG-2001.
PD
XX
           30-JAN-2001; 2001WO-US000669.
PF
XX
PR
           04-FEB-2000; 2000US-0180312P.
PR
           26-MAY-2000; 2000US-0207456P.
PR
           30-JUN-2000; 2000US-00608408.
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           03-AUG-2000; 2000US-00632366.
₽R
           21-SEP-2000; 2000US-0234687P.
           27-SEP-2000; 2000US-0236359P.
PR
           04-OCT-2000; 2000GB-00024263.
PR
XX
           (MOLE-) MOLECULAR DYNAMICS INC.
PΑ
XX
PΙ
           Penn SG, Hanzel DK, Chen W, Rank DR;
XX
           WPI; 2001-483447/52.
DR
XX
           Human genome-derived single exon nucleic acid probes useful for analyzing
PΤ
PT
           gene expression in human fetal liver.
XX
           Claim 4; SEQ ID NO 19468; 639pp + Sequence Listing; English.
PS
XX
           The invention relates to a single exon nucleic acid probe for measuring
CC
           human gene expression in a sample derived from human foetal liver. The
CC
           single exon nucleic acid probes may be used for predicting, measuring and
CC
           displaying gene expression in samples derived from human fetal liver. The
CC
           present sequence is a single exon nucleic acid probe of the invention.
CC
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CC
     Note: The sequence data for this patent did not form part of the printed
CC
     specification, but was obtained in electronic format directly from WIPO
CC
     at ftp.wipo.int/pub/published pct sequences
XX
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          65 TTGTCACTTTCCGAGGA 81
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             Db
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XX
DT
    17-OCT-2001 (first entry)
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DE
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     Probe; microarray; human; placenta; antenatal diagnosis;
KW
     genetic disorder; ss.
XX
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    Homo sapiens.
XX
PN
    WO200157272-A2.
XX
PD
    09-AUG-2001.
XX
PF
    30-JAN-2001; 2001WO-US000663.
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    26-MAY-2000; 2000US-0207456P.
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     30-JUN-2000; 2000US-00608408.
PR
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    03-AUG-2000; 2000US-00632366.
PR
    21-SEP-2000; 2000US-0234687P.
PR
    27-SEP-2000; 2000US-0236359P.
PR
    04-OCT-2000; 2000GB-00024263.
XX
PA
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
    WPI; 2001-488897/53.
XX
    Human genome-derived single exon nucleic acid probes useful for analyzing
PT
PT
    gene expression in human placenta.
XX
```

```
PS
    Claim 25; SEQ ID NO 20079; 654pp; English.
XX
CC
    The present invention relates to single exon nucleic acid probes (SENP).
    The present sequence is one such probe. The probes are useful for
CC
    producing a microarray for predicting, measuring and displaying gene
CC
    expression in samples derived from human placenta. The probes are useful
CC
     for antenatal diagnosis of human genetic disorders
CC
XX
SQ
     Sequence 180 BP; 58 A; 29 C; 41 G; 52 T; 0 U; 0 Other;
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 Best Local Similarity
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Db
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XX
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XX
DT
    06-NOV-2001 (first entry)
XX
DE
    Human bone marrow expressed single exon probe SEQ ID NO: 20005.
XX
KW
    Human; bone marrow expressed exon; gene expression analysis; probe;
KW
    microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX
os
    Homo sapiens.
XX
PN
    WO200157276-A2.
XX
PD
    09-AUG-2001.
XX
     30-JAN-2001; 2001WO-US000668.
PF
XX
PR
    04-FEB-2000; 2000US-0180312P.
    26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
    21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
₽R
     04-OCT-2000; 2000GB-00024263.
PR
XX
PΑ
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
    WPI; 2001-488900/53.
```

```
PT
    Human genome-derived single exon nucleic acid probes useful for analyzing
PT
     gene expression in human bone marrow.
XX
PS
    Example 4; SEQ ID NO 20005; 658pp + Sequence Listing; English.
XX
CC
    The present invention provides a number of single exon nucleic acid
CC
    probes which are derived from genomic sequences expressed in the human
    bone marrow. They can be used to measure gene expression in bone marrow
CC
     samples, which may enable the improved diagnosis and treatment of cancers
CC
     such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC
CC
    the probes of the invention
XX
SQ
     Sequence 180 BP; 58 A; 29 C; 41 G; 52 T; 0 U; 0 Other;
 Query Match
                          31.6%;
                                 Score 32.2; DB 4; Length 180;
                                 Pred. No. 0.28;
  Best Local Similarity
                          63.6%;
 Matches
           49; Conservative
                                 0; Mismatches
                                                 28;
                                                      Indels
                                                                 0;
                                                                     Gaps
                                                                             0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                              Db
            9 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 68
           65 TTGTCACTTTCCGAGGA 81
Qy
              11 111 111 1 111
           69 TTCTCAGTTTTCCTGGA 85
Db
RESULT 8
AAK19459
    AAK19459 standard; DNA; 180 BP.
XX
AC
    AAK19459;
XX
DT
     05-NOV-2001
                  (first entry)
XX
    Human brain expressed single exon probe SEQ ID NO: 19450.
DE
XX
    Human; brain expressed exon; gene expression analysis; probe; microarray;
KW
KW
     Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy; cancer;
KW
     ss.
XX
OS
    Homo sapiens.
XX
    WO200157275-A2.
PN
XX
PD
    09-AUG-2001.
XX
    30-JAN-2001; 2001WO-US000667.
PF
XX
     04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
```

XX

```
XX
     (MOLE-) MOLECULAR DYNAMICS INC.
PΑ
XX
PΙ
    Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
    WPI; 2001-483446/52.
XX
PΤ
    Single exon nucleic acid probes for analyzing gene expression in human
PT
    brains.
XX
PS
    Example 4; SEQ ID NO 19450; 650pp + Sequence Listing; English.
XX
CC
    The present invention provides a number of single exon nucleic acid
CC
    probes which are derived from genomic sequences expressed in the human
    brain. They can be used to measure gene expression in brain cell samples,
CC
    which may enable the diagnosis and improved treatment of nervous system
CC
CC
    diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC
    epilepsy and cancers. The present sequence is one of the probes of the
CC
    invention
XX
    Sequence 180 BP; 58 A; 29 C; 41 G; 52 T; 0 U; 0 Other;
SQ
                                Score 32.2; DB 4; Length 180;
                         31.6%;
 Best Local Similarity
                         63.6%; Pred. No. 0.28;
 Matches
           49; Conservative
                               0; Mismatches
                                                28; Indels
                                                               0; Gaps
                                                                          0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             9 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 68
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             69 TTCTCAGTTTTCCTGGA 85
Db
RESULT 9
ABS45131
ID
    ABS45131 standard; DNA; 180 BP.
XX
AC
    ABS45131;
XX
DT
    25-FEB-2003 (first entry)
XX
DE
    Human liver single exon probe, SEQ ID No 20121.
XX
KW
    Human; single exon nucleic acid probe; liver; cirrhosis;
ΚŴ
    hyperlipoproteinaemia; hyperlipidaemia; hypercholesterolaemia;
KW
    coronary heart disease; ss.
XX
OS
    Homo sapiens.
XX
    WO200157273-A2.
PN
XX
PD
    09-AUG-2001.
XX
PF
    30-JAN-2001; 2001WO-US000664.
XX
```

```
04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
PR
XΧ
     (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
                         Chen W,
                                   Rank DR;
PΙ
     Penn SG, Hanzel DK,
XX
DR
     WPI; 2001-488898/53.
XX
     Human genome-derived single exon nucleic acid probes useful for analyzing
PΤ
     gene expression in human adult liver.
PT
XX
     Claim 4; SEQ ID NO 20121; 658pp; English.
PS
XX
     The invention relates to a single exon nucleic acid probe (SENP) (I) for
CC
     measuring human gene expression in a sample derived from human adult
CC
     liver, comprising one of 13109 defined nucleotide sequences given in the
CC
CC
     specification (or complements/ fragments). The probe hybridises at high
     stringency to a nucleic acid molecule expressed in the human adult liver.
CC
     (I) may be used for predicting, measuring and displaying gene expression
CC
     in samples derived from human adult liver. The genes identified may be
CC
     involved in genetic liver diseases such as cirrhosis,
CC
     hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
CC
     associated with coronary heart disease. ABS25011-ABS51005 represent human
CC
     liver single exon nucleic acid probes of the invention. Note: The
CC
CC
     sequence information for this patent does not appear in the printed
CC
     specification but was obtained in electronic format directly from WIPO at
CC
     ftp.wipo.int/pub/published pct sequences
XX
SQ
     Sequence 180 BP; 58 A; 29 C; 41 G; 52 T; 0 U; 0 Other;
                                  Score 32.2; DB 4; Length 180;
  Query Match
                          31.6%;
                                  Pred. No. 0.28;
  Best Local Similarity
                          63.6%;
                                                                             0;
                                 0; Mismatches
                                                  28;
                                                       Indels
                                                                     Gaps
            49; Conservative
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                                            9 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 68
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
           69 TTCTCAGTTTTCCTGGA 85
RESULT 10
ABS19713
     ABS19713 standard; DNA; 180 BP.
ID
XX
AC
     ABS19713;
XX
DT
     19-AUG-2002 (first entry)
XX
```

Human genome-derived single exon probe ORF from lung SEQ ID No 19704. DΕ XX Human; ds; single exon probe; asthma; lung cancer; COPD; ILD; KW chronic obstructive pulmonary disease; interstitial lung disease; KW familial idiopathic pulmonary fibrosis; neurofibromatosis; ΚW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease; KW KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis; pulmonary histiocytosis; lymphangioleiomyomtosis; Karagener syndrome; KW KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia; KW primary ciliary dyskinesis; pulmonary hypertension; hyaline membrane disease; open reading frame; ORF. KW XX OS Homo sapiens. XXWO200186003-A2. PNXX PD15-NOV-2001. XX 30-JAN-2001; 2001WO-US000665. PFXX 04-FEB-2000; 2000US-0180312P. PR 26-MAY-2000; 2000US-0207456P. PR 30-JUN-2000; 2000US-00608408. PR 03-AUG-2000; 2000US-00632366. PR 21-SEP-2000; 2000US-0234687P. PR 27-SEP-2000; 2000US-0236359P. PR PR 04-OCT-2000; 2000GB-00024263. XX (MOLE-) MOLECULAR DYNAMICS INC. PA XX PΙ Hanzel DK, Chen W, Rank DR; Penn SG, XX DR WPI; 2002-114183/15. XX PT Spatially-addressable set of single exon nucleic acid probes, used to PTmeasure gene expression in human lung samples. XX PS Claim 4; SEQ ID NO 19704; 634pp; English. XX The invention relates to a spatially-addressable set of single exon CC CC nucleic acid probes for measuring gene expression in a sample derived from human lung comprising single exon nucleic acid probes having one of CC CC 12614 nucleic acid sequences mentioned in the specification, or their complements or the 12387 open reading frames derived from the 12614CC probes. Also included are a microarray comprising the novel set of probes CC ; the novel set of probes which hybridise at high stringency to a nucleic CC CC acid expressed in the human lung; measuring gene expression in a sample derived from human lung, comprising (a) contacting the array with a CC collection of detectably labeled nucleic acids derived from human lung CC mRNA, and (b) measuring the label detectably bound to each probe of the CC CC array; identifying exons in a eukaryotic genome, comprising (a) CC algorithmically predicting at least one exon from genomic sequences of the eukaryote; and (b) detecting specific hybridisation of detectably CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe, CC CC having a fragment identical to the predicted exon, the probe is included CC in the above mentioned microarray; assigning exons to a single gene, CC comprising (a) identifying exons from genomic sequence by the method

```
CC
     above and (b) measuring the expression of each of the exons in several
CC
     tissues and/or cell types using hybridisation to a single exon
     microarrays having a probe with the exon, where a common pattern of
CC
CC
     expression of the exons in the tissues and/or cell types indicates that
     the exons should be assigned to a single gene; a peptide comprising one
CC
CC
     of 12011 sequences, mentioned in the specification, or encoded by the
CC
     probes/open reading frames (ORF). The probes are used for gene expression
CC
     analysis, and for identifying exons in a gene, particularly using human
CC
     lung derived mRNA and for the study of lung diseases such as asthma, lung
CC
     cancer, chronic obstructive pulmonary disease (COPD), interstitial lung
CC
     disease (ILD), familial idiopathic pulmonary fibrosis, neurofibromatosis,
CC
     tuberous sclerosis, Gaucher's disease, Niemann-Pick disease, Hermansky-
CC
     Pudlak syndrome, sarcoidosis, pulmonary haemosiderosis, pulmonary
CC
     histiocytosis, lymphangioleiomyomtosis, pulmonary alveolar proteinosis,
CC
     Karagener syndrome, fibrocystic pulmonary dysplasia, primary ciliary
CC
     dyskinesis, pulmonary hypertension and hyaline membrane disease. The
CC
     present sequence is a single exon probe open reading frame of the
     invention. Note: The sequence data for this patent did not form part of
CC
CC
     the printed specification, but was obtained in electronic format directly
CC
     from WIPO at ftp.wipo.int/pub/published pct sequences
XX
SQ
     Sequence 180 BP; 58 A; 29 C; 41 G; 52 T; 0 U; 0 Other;
  Query Match
                          31.6%;
                                 Score 32.2; DB 6; Length 180;
  Best Local Similarity
                          63.6%;
                                 Pred. No. 0.28;
  Matches
           49; Conservative
                                 0; Mismatches
                                                 28;
                                                      Indels
                                                                    Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
              9 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 68
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 11 11 1 11
Db
           69 TTCTCAGTTTTCCTGGA 85
RESULT 11
ABA58823
    ABA58823 standard; DNA; 494 BP.
XX
AC
    ABA58823;
XX
DT
     01-FEB-2002 (first entry)
XX
DE
     Human foetal liver single exon nucleic acid probe #7128.
XX
KW
     Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
XX
OS
     Homo sapiens.
XX
PN
    WO200157277-A2.
XX
PD
     09-AUG-2001.
XX
PF
     30-JAN-2001; 2001WO-US000669.
XX
     04-FEB-2000; 2000US-0180312P.
PR
```

```
26-MAY-2000; 2000US-0207456P.
PR
    30-JUN-2000; 2000US-00608408.
PR
    03-AUG-2000; 2000US-00632366.
PR
    21-SEP-2000; 2000US-0234687P.
PR
    27-SEP-2000; 2000US-0236359P.
PR
    04-OCT-2000; 2000GB-00024263.
PR
XX
     (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
    Penn SG, Hanzel DK, Chen W, Rank DR;
PΙ
XX
    WPI; 2001-483447/52.
DR
XX
    Human genome-derived single exon nucleic acid probes useful for analyzing
РT
PT
    gene expression in human fetal liver.
XX
    Claim 1; SEQ ID NO 7128; 639pp + Sequence Listing; English.
PS
XX
    The invention relates to a single exon nucleic acid probe for measuring
CC
    human gene expression in a sample derived from human foetal liver. The
CC
    single exon nucleic acid probes may be used for predicting, measuring and
CC
    displaying gene expression in samples derived from human fetal liver. The
CC
    present sequence is a single exon nucleic acid probe of the invention.
CC
    Note: The sequence data for this patent did not form part of the printed
CC
     specification, but was obtained in electronic format directly from WIPO
CC
CC
     at ftp.wipo.int/pub/published pct sequences
XX
    Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 4; Length 494;
  Ouerv Match
  Best Local Similarity 63.6%; Pred. No. 0.38;
           49; Conservative
                                0; Mismatches
                                                28; Indels
                                                               0; Gaps
                                                                           0;
 Matches
Qу
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
             280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
Db
Qy
           65 TTGTCACTTTCCGAGGA 81
             Db
          340 TTCTCAGTTTTCCTGGA 356
RESULT 12
AAI38528
    AAI38528 standard; DNA; 494 BP.
ΙD
XX
AC
    AAI38528;
XX
DT
     17-OCT-2001 (first entry)
XX
DE
     Probe #7214 used to measure gene expression in human placenta sample.
XX
KW
     Probe; microarray; human; placenta; antenatal diagnosis;
ΚW
     genetic disorder; ss.
XX
OS
     Homo sapiens.
XX
```

```
PN
    WO200157272-A2.
XX
PD
     09-AUG-2001.
XX
PF
    30-JAN-2001; 2001WO-US000663.
XX
     04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
XX
PΑ
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
    WPI; 2001-488897/53.
XX
    Human genome-derived single exon nucleic acid probes useful for analyzing
PT
PT
     gene expression in human placenta.
XX
PS
    Claim 25; SEQ ID NO 7214; 654pp; English.
XX
     The present invention relates to single exon nucleic acid probes (SENP).
CC
     The present sequence is one such probe. The probes are useful for
CC
    producing a microarray for predicting, measuring and displaying gene
CC
     expression in samples derived from human placenta. The probes are useful
CC
     for antenatal diagnosis of human genetic disorders
CC
XX
     Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;
SO
                         31.6%; Score 32.2; DB 4; Length 494;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.38;
                                0; Mismatches 28; Indels
           49; Conservative
                                                                0; Gaps
                                                                            0;
  Matches
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                             Db
         280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
          65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
         340 TTCTCAGTTTTCCTGGA 356
RESULT 13
AAK32713
ID
    AAK32713 standard; DNA; 494 BP.
XX
AC
    AAK32713;
XX
DT
     06-NOV-2001 (first entry)
XX
     Human bone marrow expressed single exon probe SEQ ID NO: 7270.
DE
XX
KW
     Human; bone marrow expressed exon; gene expression analysis; probe;
```

```
KW
    microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX
OS
     Homo sapiens.
XX
ΡN
    WO200157276-A2.
XX
PD
     09-AUG-2001.
XX
PF
     30-JAN-2001; 2001WO-US000668.
XX
PR
     04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
     04-OCT-2000; 2000GB-00024263.
PR
XX
PΑ
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
    WPI; 2001-488900/53.
XX
PT
    Human genome-derived single exon nucleic acid probes useful for analyzing
     gene expression in human bone marrow.
PT
XX
PS
     Example 4; SEQ ID NO 7270; 658pp + Sequence Listing; English.
XX
CC
     The present invention provides a number of single exon nucleic acid
     probes which are derived from genomic sequences expressed in the human
CC
CC
     bone marrow. They can be used to measure gene expression in bone marrow
     samples, which may enable the improved diagnosis and treatment of cancers
CC
     such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC
     the probes of the invention
CC
XX
     Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;
SQ
                                 Score 32.2; DB 4; Length 494;
  Query Match
                         31.6%;
  Best Local Similarity
                         63.6%; Pred. No. 0.38;
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                    Gaps
                                                                            0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                             Db
         280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
          65 TTGTCACTTTCCGAGGA 81
Qу
              340 TTCTCAGTTTTCCTGGA 356
RESULT 14
AAK06977
    AAK06977 standard; DNA; 494 BP.
ID
XX
AC
    AAK06977:
XX
```

```
DT
     05-NOV-2001 (first entry)
XX
DΕ
     Human brain expressed single exon probe SEQ ID NO: 6968.
XX
ΚW
    Human; brain expressed exon; gene expression analysis; probe; microarray;
KW
    Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy; cancer;
KW
XX
OS
    Homo sapiens.
XX
PN
    WO200157275-A2.
XX
PD
    09-AUG-2001.
XX
PF
    30-JAN-2001; 2001WO-US000667.
XX
     04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
PR
XX
     (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
PI
     Penn SG, Hanzel DK, Chen W,
                                   Rank DR;
XX
DR
    WPI; 2001-483446/52.
XX
     Single exon nucleic acid probes for analyzing gene expression in human
PT
PT
    brains.
XX
PS
    Example 4; SEQ ID NO 6968; 650pp + Sequence Listing; English.
XX
CC
    The present invention provides a number of single exon nucleic acid
CC
     probes which are derived from genomic sequences expressed in the human
CC
    brain. They can be used to measure gene expression in brain cell samples,
    which may enable the diagnosis and improved treatment of nervous system
CC
CC
    diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC
    epilepsy and cancers. The present sequence is one of the probes of the
CC
    invention
XX
SO
     Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;
  Query Match
                          31.6%;
                                 Score 32.2; DB 4; Length 494;
  Best Local Similarity
                          63.6%;
                                 Pred. No. 0.38;
  Matches
           49; Conservative
                                 0; Mismatches
                                                  28;
                                                      Indels
                                                                 0;
                                                                    Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                             Db
          280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
           65 TTGTCACTTTCCGAGGA 81
Qγ
              11 111 111 1 111
Db
          340 TTCTCAGTTTTCCTGGA 356
```

```
RESULT 15
ABS32432
    ABS32432 standard; DNA; 494 BP.
ID
XX
AC
    ABS32432;
XX
DT
    25-FEB-2003 (first entry)
XX
DE
    Human liver single exon probe, SEQ ID No 7422.
XX
     Human; single exon nucleic acid probe; liver; cirrhosis;
KW
     hyperlipoproteinaemia; hyperlipidaemia; hypercholesterolaemia;
KW
     coronary heart disease; ss.
KW
XX
OS
     Homo sapiens.
XX
PN
     WO200157273-A2.
XX
PD
     09-AUG-2001.
XX
     30-JAN-2001; 2001WO-US000664.
PF
XX
     04-FEB-2000; 2000US-0180312P.
PR
     26-MAY-2000; 2000US-0207456P.
PR
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
XX
PΑ
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR
     WPI; 2001-488898/53.
XX
     Human genome-derived single exon nucleic acid probes useful for analyzing
PT
PΤ
     gene expression in human adult liver.
XX
PS
     Claim 1; SEQ ID NO 7422; 658pp; English.
XX
     The invention relates to a single exon nucleic acid probe (SENP) (I) for
CC
     measuring human gene expression in a sample derived from human adult
CC
     liver, comprising one of 13109 defined nucleotide sequences given in the
CC
     specification (or complements/ fragments). The probe hybridises at high
CC
     stringency to a nucleic acid molecule expressed in the human adult liver.
CC
CC
     (I) may be used for predicting, measuring and displaying gene expression
CC
     in samples derived from human adult liver. The genes identified may be
CC
     involved in genetic liver diseases such as cirrhosis,
CC
     hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
     associated with coronary heart disease. ABS25011-ABS51005 represent human
CC
CC
     liver single exon nucleic acid probes of the invention. Note: The
CC
     sequence information for this patent does not appear in the printed
CC
     specification but was obtained in electronic format directly from WIPO at
CC
     ftp.wipo.int/pub/published pct sequences
XX
```

```
Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;
SQ
  Query Match
                          31.6%;
                                 Score 32.2; DB 4; Length 494;
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.38;
  Matches
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                0; Gaps
                                                                            0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qγ
                                            280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
Db
         340 TTCTCAGTTTTCCTGGA 356
RESULT 16
ABS07509
ID
    ABS07509 standard; DNA; 494 BP.
XX
AC
    ABS07509;
XX
DT
     19-AUG-2002 (first entry)
XX
    Human genome-derived single exon probe from lung SEQ ID No 7500.
DE
XX
     Human; ds; single exon probe; asthma; lung cancer; COPD; ILD;
KW
     chronic obstructive pulmonary disease; interstitial lung disease;
KW
     familial idiopathic pulmonary fibrosis; neurofibromatosis;
KW
     tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
KW
     Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
KW
     pulmonary histiocytosis; lymphangioleiomyomtosis; Karagener syndrome;
KW
KW
     pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
     primary ciliary dyskinesis; pulmonary hypertension;
KW
KW
     hyaline membrane disease.
XX
OS
     Homo sapiens.
XX
PN
    W0200186003-A2.
XX
PD
    15-NOV-2001.
XX
PF
     30-JAN-2001; 2001WO-US000665.
XX
PR
     04-FEB-2000; 2000US-0180312P.
     26-MAY-2000; 2000US-0207456P.
PR
     30-JUN-2000; 2000US-00608408.
PR
     03-AUG-2000; 2000US-00632366.
PR
     21-SEP-2000; 2000US-0234687P.
PR
     27-SEP-2000; 2000US-0236359P.
PR
     04-OCT-2000; 2000GB-00024263.
PR
XX
PΑ
     (MOLE-) MOLECULAR DYNAMICS INC.
XX
PΙ
     Penn SG, Hanzel DK, Chen W,
                                   Rank DR;
XX
DR
     WPI; 2002-114183/15.
```

Spatially-addressable set of single exon nucleic acid probes, used to measure gene expression in human lung samples.

PT XX PS

PT

Claim 1; SEQ ID NO 7500; 634pp; English.

XX CC

CC

СĊ

CC

CC CC

CC

CC

CC

CC

The invention relates to a spatially-addressable set of single exon nucleic acid probes for measuring gene expression in a sample derived from human lung comprising single exon nucleic acid probes having one of 12614 nucleic acid sequences mentioned in the specification, or their complements or the 12387 open reading frames derived from the 12614 probes. Also included are a microarray comprising the novel set of probes ; the novel set of probes which hybridise at high stringency to a nucleic acid expressed in the human lung; measuring gene expression in a sample derived from human lung, comprising (a) contacting the array with a collection of detectably labeled nucleic acids derived from human lung mRNA, and (b) measuring the label detectably bound to each probe of the array; identifying exons in a eukaryotic genome, comprising (a) algorithmically predicting at least one exon from genomic sequences of the eukaryote; and (b) detecting specific hybridisation of detectably labeled nucleic acids from eukaryote lung mRNA, to a single exon probe, having a fragment identical to the predicted exon, the probe is included in the above mentioned microarray; assigning exons to a single gene, comprising (a) identifying exons from genomic sequence by the method above and (b) measuring the expression of each of the exons in several tissues and/or cell types using hybridisation to a single exon microarrays having a probe with the exon, where a common pattern of expression of the exons in the tissues and/or cell types indicates that the exons should be assigned to a single gene; a peptide comprising one of 12011 sequences, mentioned in the specification, or encoded by the probes/open reading frames (ORF). The probes are used for gene expression analysis, and for identifying exons in a gene, particularly using human lung derived mRNA and for the study of lung diseases such as asthma, lung cancer, chronic obstructive pulmonary disease (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease, Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomtosis, pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic pulmonary dysplasia, primary ciliary dyskinesis, pulmonary hypertension and hyaline membrane disease. The present sequence is a single exon probe of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published pct sequences

CC XX SO

Sequence 494 BP; 155 A; 81 C; 92 G; 166 T; 0 U; 0 Other;

```
Query Match
                     31.6%; Score 32.2; DB 6; Length 494;
 Best Local Similarity
                     63.6%;
                            Pred. No. 0.38;
 Matches
          49; Conservative
                           0; Mismatches
                                          28;
                                                                0;
                                              Indels
                                                          Gaps
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                         Db
        280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
```

```
Qy 65 TTGTCACTTTCCGAGGA 81
|| || || || || || Db 340 TTCTCAGTTTTCCTGGA 356
```

```
RESULT 17
AAZ99413
     AAZ99413 standard; DNA; 500 BP.
XX
    AAZ99413;
AC
XX
DT
     03-JUL-2000 (first entry)
XX
DE
     Trans-spliced product of the CFTR target pre-mRNA and a PTM.
XX
KW
     Pre-mRNA molecule; gene repair; pre-trans-splicing molecule;
KW
     gene regulation; targeted cell death;
ΚW
     cystic fibrosis trans-membrane regulator gene; ss.
XX
OS
     Homo sapiens.
XX
PN
     WO200009734-A2.
XX
PD
     24-FEB-2000.
XX
PF
     12-AUG-1999;
                    99WO-US018371.
XX
PR
     13-AUG-1998;
                    98US-00133717.
PΈ
     23-SEP-1998;
                    98US-00158863.
XX
     (INTR-) INTRONN HOLDINGS LLC.
PA
XX
PΙ
     Mitchell LG, Garcia-Blanco MA;
XX
DR '
     WPI: 2000-224360/19.
XX
PT
     Novel pre-trans-splicing molecules for use in gene regulation, gene
     repair and targeted cell death particularly gene repair of cystic
PT
PT
     fibrosis trans-membrane regulator gene.
XX
PS
     Example 8; Fig 15; 79pp; English.
XX
CC
     The specification describes a pre-trans-splicing molecule (PTM) which
     contains one or more target binding domains, a 3' splice region
CC
CC
     comprising a branch point, a pyrimidine tract and a 3' splice acceptor
     site, a spacer region separating the mRNA splice region from the target
CC
CC
     binding domain, and a nucleotide sequence to be trans-spliced. The method
     is used for the in vivo production of a trans-spliced molecule in a
CC
     subset of cells. The PTM is used for producing chimeric mRNA molecule by
CC
     contacting it with target pre mRNA which is useful for gene regulation,
CC
CC
     gene repair and targeted cell death particularly repair of cystic
CC
     fibrosis trans-membrane regulator (CFTR) gene. The present sequence
CC
     represents the trans-spliced product of the CFTR target pre-mRNA and a
CC
     PTM of the invention
XX
     Sequence 500 BP; 125 A; 127 C; 102 G; 146 T; 0 U; 0 Other;
SO
                                  Score 32.2; DB 3; Length 500;
                          31.6%;
                          63.6%; Pred. No. 0.38;
  Best Local Similarity
                                 0; Mismatches
                                                  28; Indels
                                                                              0;
  Matches
           49; Conservative
                                                                  0; Gaps
```

```
5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              Db
          128 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 187
           65 TTGTCACTTTCCGAGGA 81
Qу
             Db
          188 TTCTCAGTTTTCCTGGA 204
RESULT 18
AB073502
ID
     ABQ73502 standard; DNA; 500 BP.
XX
AC
     ABQ73502;
XX
DT
     02-OCT-2002 (first entry)
XX
DE
     Pre-trans-splicing molecule related oligonucleotide #9.
XX
KW
     Pre-trans-splicing molecule; PTM; spliceosome; cytostatic; gene therapy;
KW
     immunosuppressive; antimicrobial; gene regulation; gene repair; cancer;
KW
     targeted cell death; genetic disorder; infectious disorder;
KW
     autoimmune disease; proliferative disorder; PCR primer; ss.
XX
     Synthetic.
OS
XX
     WO200253581-A2.
PN
XX
PD
     11-JUL-2002.
XX
     08-JAN-2002; 2002WO-US000416.
PF
XX
PR
     08-JAN-2001; 2001US-00756095.
     08-JAN-2001; 2001US-00756096.
PR
     08-JAN-2001; 2001US-00756097.
PR
     20-APR-2001; 2001US-00838858.
PR
     29-AUG-2001; 2001US-00941492.
PR
XX
PA
     (INTR-) INTRONN INC.
XX
PΙ
    Mitchell LG, Garcia-Blanco MA, Baker CC, Puttaraju M;
ΡI
    Mansfield GS, Chao H;
XX
DR
    WPI; 2002-566693/60.
XX
PT
    Novel cell having pre-trans-splicing molecules with target binding
     domains that target binding of PTM to pre-mRNA, 3' or 5' splice region,
PT
PT
     spacer region, nucleotide sequence to be trans-spliced to target-pre-
PT
    mRNA.
XX
PS
     Example; Fig 15A-B; 229pp; English.
XX
    The present invention describes a cell (I) comprising pre-trans-splicing
CC
CC
    molecules (PTMs) (II) which have one or more target binding domains (IIa)
CC
     that target binding of PTM to pre-mRNA, 3' splice region (IIb) that
CC
     includes branch point pyrimidine tract and 3'splice acceptor site, or 5'
```

```
CC
     splice site (IIc), spacer region (IId) that separates RNA splice site
CC
     from target binding domain, and nucleotide sequence to (IIe) be trans-
CC
     spliced to target-pre-mRNA. Optionally, the cell comprises (II) either
CC
     comprising: (A) (IIb) and (IIe); or (B) (IIc), (IId) and (IIe). The cell
     may comprise a recombinant vector expressing (II). (I) has cytostatic,
CC
CC
     immunosuppressive and antimicrobial activities, and can be used in gene
CC
     therapy. (II) comprising one or more (preferably two or more) (IIa) and
CC
     (IIb) (or (IIc)), (IId) and (IIe), or (II) comprising either (A) or (B)
CC
     (excluding (IId)), is useful for producing a chimeric RNA molecule in a
CC
     cell which involves contacting a target pre-mRNA expressed in the cell
     with (II) that is recognised by nuclear splicing components. The chimeric
CC
CC
     RNA produced comprises sequences encoding a toxin or translatable
CC
     protein. The nucleotide sequence to be trans-spliced to target pre-mRNA
CC
     preferably comprises nucleotide sequences comprising exons 1-10 of cystic
CC
     fibrosis trans-membrane conductance regulator (CFTR). The chimeric RNA
CC
     molecule produced using (II) which either comprises (A) or (B) further
CC
     comprises a nucleotide sequence tag. (I) can be used for gene regulation,
CC
     gene repair and targeted cell death. (I) can be used for the treatment of
CC
     various diseases including genetic, infectious or autoimmune diseases and
CC
     proliferative disorders such as cancer and to regulate gene expression in
CC
     plants. ABQ73414 to ABQ73536 represent sequences used in the
CC
     exemplification of the present invention
XX
SO
     Sequence 500 BP; 125 A; 128 C; 101 G; 146 T; 0 U; 0 Other;
  Query Match
                          31.6%;
                                 Score 32.2; DB 6; Length 500;
  Best Local Similarity
                          63.6%;
                                 Pred. No. 0.38;
            49; Conservative
                                 0; Mismatches
                                                 28;
                                                      Indels
                                                                            0:
                                                                    Gaps
QУ
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
                              128 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 187
Db
Qу
           65 TTGTCACTTTCCGAGGA 81
              Db
          188 TTCTCAGTTTTCCTGGA 204
RESULT 19
ABZ24468
     ABZ24468 standard; DNA; 795 BP.
XX
AC
     ABZ24468;
XX
DT
     21-MAR-2003
                 (first entry)
XX
DΕ
     Cystic fibrosis transmembrane conductance regulator gene exon 10.
XX
     Cystic fibrosis transmembrane conductance regulator; CFTR; human;
KW
KW
     cystic fibrosis; nucleic acid detection; quality assurance; validation;
KW
     diagnosis; ds.
XX
os
     Homo sapiens.
XX
PN
     WO200296925-A1.
XX
PD
     05-DEC-2002.
```

```
XX
ΡF
     24-MAY-2002; 2002WO-US016504.
XX
PR
     25-MAY-2001; 2001US-00866293.
XX
PA
     (MAIN-) MAINE MEDICAL CENT RES INST.
PΑ
     (MAIN-) MAINE MOLECULAR QUALITY CONTROLS INC.
XX
PΙ
     Gordon J, Rundell CA;
XX
DR
    WPI; 2003-140437/13.
XX
PT
     Control DNA constructs useful in nucleic acid assays, has vector portion
PT
     for expression in a cell and a target nucleic acid comprising fragments
PT
     which specify component associated with disease state or environmental
РΤ
     condition.
XX
PS
     Disclosure; Page 74-75; 76pp; English.
XX
    The present sequence is the nucleotide sequence of exon 10 of the human
CC
CC
     cystic fibrosis transmembrane conductance regulator (CFTR) gene. Many of
     the most common disease-causing mutations are in exon 10 and exon 11 (see
CC
CC
     ABZ24469) of the CFTR gene, and genetic screening for these mutations is
     therefore advantageous for early diagnosis of cystic fibrosis. The
CC
CC
     invention provides control DNA constructs useful in nucleic acid assays.
     The DNA constructs have a vector portion for expression in a cell and a
CC
     target nucleic acid comprising 2 or more nucleic acid fragments, where
CC
CC
     each fragment specifies a component associated with a disease state, an
CC
     environmental condition or a biological organism. Each fragment may
     comprise at least 1 exon of a gene, and is especially a CFTR exon,
CC
     particularly exon 10 and exon 11. The DNA constructs provide controls
CC
CC
     useful for quality assurance in the diagnostic detection of complex
     genetic diseases such as cystic fibrosis, and for quality assurance in
CC
CC
     nucleic acid assays to detect components associated with an environmental
CC
     condition or a biological organism
XX
SO
     Sequence 795 BP; 251 A; 143 C; 135 G; 266 T; 0 U; 0 Other;
                                 Score 32.2; DB 7; Length 795;
  Query Match
                         31.6%;
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.44;
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                    Gaps
                                                                            0:
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                             Db
          369 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 428
           65 TTGTCACTTTCCGAGGA 81
Qу
              429 TTCTCAGTTTTCCTGGA 445
Db
RESULT 20
     ADE77694 standard; DNA; 831 BP.
ID
XX
AC
    ADE77694:
XX
```

```
DT
     29-JAN-2004 (first entry)
XX
DE
     Human cystic fibrosis conductance transmembrane regulator exon 10 DNA.
XX
     ds; human; CFTR; human leukocyte antigen; HLA; genetic testing;
KW
    carrier screening; genotyping; profiling; polymorphic;
KW
     multiplexed elongation assay; enzymatic recognition;
KW
KW
     cystic fibrosis conductance transmembrane regulator;
KW
     single nucleotide polymorphism; SNP.
XX
OS
     Homo sapiens.
XX
ΡN
     WO2003034029-A2.
XX
PD
     24-APR-2003.
XX
PF
     15-OCT-2002; 2002WO-US033012.
XX
PR
     15-OCT-2001; 2001US-0329427P.
     15-OCT-2001; 2001US-0329428P.
PR
PR
     15-OCT-2001; 2001US-0329619P.
PR
     15-OCT-2001; 2001US-0329620P.
     14-MAR-2002; 2002US-0364416P.
PR
XX
PA
     (BIOA-) BIOARRAY SOLUTIONS LTD.
XX
PΙ
     Li AX, Hashmi G, Seul M;
XX
DR
     WPI; 2003-393553/37.
XX
PΤ
     Concurrent interrogation of a number of polymorphic sites, useful for
PT
     genetic testing, carrier screening, genetic profiling, and identity
PT
     testing, comprises conducting a multiplexed elongation assay using
PT
     probes.
XX
PS
     Example 12; Page 54; 143pp; English.
XX
CC
     This invention relates to a novel method for the concurrent interrogation
CC
     of a number of polymorphic sites in the presence of, and without
CC
     interference from, non-designated polymorphic sites. Specifically, it
CC
     comprises conducting a multiplexed elongation assay by applying one or
CC
     more temperature cycles to achieve linear amplification of the target or
CC
     a combination of annealing and elongation steps under temperature-
CC
     controlled conditions. Furthermore, this detection method uses probe
CC
     extension or elongation and relies on enzymatic recognition, a superior
CC
     technique that no longer depends on differential hybridisation. The
```

genotyping or genetic profiling, and identity testing. This polynucleotide is the human cystic fibrosis conductance transmembrane regulator (CFTR) exon 10 DNA sequence containing single nucleotide polymorphisms, used in an exemplification of the invention.

transmembrane regulator (CFTR) or the human leukocyte antigen (HLA)

polymorphic sites is useful for genetic testing, carrier screening,

genes. In addition, concurrent interrogation of a multiplicity of

can identify mutations within the cystic fibrosis conductance

present invention describes probes and methods useful for identifying or

detecting polymorphisms at one or more designated sites, such that they

CC

```
Sequence 831 BP; 263 A; 140 C; 141 G; 287 T; 0 U; 0 Other;
SO
  Query Match
                         31.6%; Score 32.2; DB 9; Length 831;
  Best Local Similarity
                         63.6%; Pred. No. 0.44;
 Matches
          49; Conservative
                                0; Mismatches
                                                28; Indels
                                                               0; Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
             Db
         328 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 387
          65 TTGTCACTTTCCGAGGA 81
Qy
             Db
         388 TTCTCAGTTTTCCTGGA 404
RESULT 21
AAT04005
ΙD
    AAT04005 standard; cDNA; 2640 BP.
XX
AC
    AAT04005;
XX
DT
    25-MAR-2003 (revised)
DT
    02-MAY-1996 (first entry)
XX
DΕ
    Truncated cystic fibrosis transmembrane conductance regulator cDNA.
XX
KW
    Cystic fibrosis; transmembrane conductance; N-terminal; soluble;
     truncated; chloride ion channel; gene therapy; CFTR; regulator;
KW
KW
    epithelial cells; anion; recombinant production; ss.
XX
    Homo sapiens.
OS
XX
FΗ
                    Location/Qualifiers
    Key
FT
                    133. .2640
    CDS
FT
                    /*tag= a
                    /note= "truncated N-terminal CFTR protein"
FT
XX
PN
    WO9525796-A1.
ΧX
PD
    28-SEP-1995.
XX
PF
    23-MAR-1995;
                   95WO-US003680.
XX
PR
    23-MAR-1994;
                   94US-00216971.
XX
PΑ
    (IOWA ) UNIV IOWA STATE RES FOUND INC.
XX
PI
    Welsh MJ, Sheppard DM;
XX
DR
    WPI; 1995-344617/44.
DR
    P-PSDB; AAR79835.
XX
PT
    New truncated CFTR polypeptide - functions as a regulated epithelial cell
PT
XX '
    anion channel, used for treating cystic fibrosis.
PS
    Claim 5; Page 67-70; 85pp; English.
XX
```

```
CC
     AAT04005 encodes AAR79835 a truncated N-terminal portion of the cystic
CC
     fibrosis transmembrane conductance regulator (CFTR), which can be used to
CC
     regulate the opening and closing of epithelial cell anion (chloride ion)
CC
     channels. The truncated cDNA is useful in CF gene therapy, as it is more
CC
     readily accommodated by available gene therapy vectors, and more easily
     expressed than full length CFTR. The expressed truncated CFTR protein may
CC
CC
     be more soluble and therefore more readily purified from host cells,
CC
     useful in the recombinant prodn. of CFTR. (Updated on 25-MAR-2003 to
CC
     correct PI field.)
XX
SQ
     Sequence 2640 BP; 836 A; 509 C; 584 G; 711 T; 0 U; 0 Other;
  Query Match
                          31.6%; Score 32.2; DB 2; Length 2640;
  Best Local Similarity
                          63.6%; Pred. No. 0.64;
            49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                0; Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
              Db
         1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Qy
           65 TTGTCACTTTCCGAGGA 81
              11 111 111 1 111
Db
         1605 TTCTCAGTTTTCCTGGA 1621
RESULT 22
ABO73521
     ABQ73521 standard; DNA; 3069 BP.
XX
AC
     ABQ73521;
XX
DT
     02-OCT-2002 (first entry)
XX
DE
    Mouse factor VIII PTM nucleotide sequence.
XX
KW
     Pre-trans-splicing molecule; PTM; spliceosome; cytostatic; gene therapy;
KW
     immunosuppressive; antimicrobial; gene regulation; gene repair; cancer;
KW
     targeted cell death; genetic disorder; infectious disorder;
KW
     autoimmune disease; proliferative disorder; gene; ds.
XX
OS
    Mus sp.
OS
    Synthetic.
XX
PN
    WO200253581-A2.
XX
PD
    11~JUL-2002.
XX
PF
    08-JAN-2002; 2002WO-US000416.
XX
PR
    08-JAN-2001; 2001US-00756095.
PR
    08-JAN-2001; 2001US-00756096.
     08-JAN-2001; 2001US-00756097.
PR
PR
    20-APR-2001; 2001US-00838858.
PR
    29-AUG-2001; 2001US-00941492.
XX
PA
     (INTR-) INTRONN INC.
XX
```

```
PΙ
     Mitchell LG,
                   Garcia-Blanco MA, Baker CC, Puttaraju M;
PΙ
     Mansfield GS,
                    Chao H;
XX
DR
     WPI; 2002-566693/60.
XX
     Novel cell having pre-trans-splicing molecules with target binding
PT
PT
     domains that target binding of PTM to pre-mRNA, 3' or 5' splice region,
PT
     spacer region, nucleotide sequence to be trans-spliced to target-pre-
PT
     mRNA.
XX
PS
     Example; Fig 43B; 229pp; English.
XX
     The present invention describes a cell (I) comprising pre-trans-splicing
CC
     molecules (PTMs) (II) which have one or more target binding domains (IIa)
CC
     that target binding of PTM to pre-mRNA, 3' splice region (IIb) that
CC
CC
     includes branch point pyrimidine tract and 3'splice acceptor site, or 5'
     splice site (IIc), spacer region (IId) that separates RNA splice site
CC
CC
     from target binding domain, and nucleotide sequence to (IIe) be trans-
     spliced to target-pre-mRNA. Optionally, the cell comprises (II) either comprising: (A) (IIb) and (IIe); or (B) (IIc), (IId) and (IIe). The cell
CC
CC
CC
     may comprise a recombinant vector expressing (II). (I) has cytostatic,
CC
     immunosuppressive and antimicrobial activities, and can be used in gene
CC
     therapy. (II) comprising one or more (preferably two or more) (IIa) and
CC
     (IIb) (or (IIc)), (IId) and (IIe), or (II) comprising either (A) or (B)
CC
     (excluding (IId)), is useful for producing a chimeric RNA molecule in a
CC
     cell which involves contacting a target pre-mRNA expressed in the cell
CC
     with (II) that is recognised by nuclear splicing components. The chimeric
CC
     RNA produced comprises sequences encoding a toxin or translatable
CC
     protein. The nucleotide sequence to be trans-spliced to target pre-mRNA
CC
     preferably comprises nucleotide sequences comprising exons 1-10 of cystic
CC
     fibrosis trans-membrane conductance regulator (CFTR). The chimeric RNA
CC
     molecule produced using (II) which either comprises (A) or (B) further
CC
     comprises a nucleotide sequence tag. (I) can be used for gene regulation,
     gene repair and targeted cell death. (I) can be used for the treatment of
CC
CC
     various diseases including genetic, infectious or autoimmune diseases and
CC
     proliferative disorders such as cancer and to regulate gene expression in
CC
     plants. ABQ73414 to ABQ73536 represent sequences used in the
CC
     exemplification of the present invention
XX
SO
     Sequence 3069 BP; 955 A; 609 C; 662 G; 843 T; 0 U; 0 Other;
                                  Score 32.2; DB 6; Length 3069;
                          31.6%;
  Best Local Similarity
                          63.6%;
                                  Pred. No. 0.67;
 Matches
            49; Conservative
                                 0; Mismatches
                                                   28;
                                                        Indels
                                                                  0; Gaps
                                                                               0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                              Db
           21 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 80
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
Db
           81 TTCTCAGTTTTCCTGGA 97
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RESULT 23 AAF84742

ID AAF84742 standard; DNA; 4443 BP.

```
XX
AC
     AAF84742;
XX
DT
     29-JUN-2001 (first entry)
XX
DE
     DNA encoding cystic fibrosis transmembrane conductance regulator (CFTR).
XX
KW
     Cystic fibrosis transmembrane conductance regulator; CFTR;
KW
     cystic fibrosis; CTFR dimerisation; ss.
XX
OS
     Homo sapiens.
XX
FH
     Key
                     Location/Qualifiers
                     1. .4443
FT
     CDS
FT
                     /*tag= a
FT
                     /transl except= (pos: 2497. .2499, aa: Leu)
                     /product= "cystic fibrosis transmembrane conductance
FT
FT
                     regulator (CFTR)"
XX
ΡN
     WO200125421-A2.
XX
PD
     12-APR-2001.
XX
PF
     06-OCT-2000; 2000WO-US027900.
XX
PR
     06-OCT-1999;
                    99US-0157996P.
     11-FEB-2000; 2000US-0181892P.
PR
     14-FEB-2000; 2000US-0182373P.
PR
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
DR
     WPI; 2001-273576/28.
     P-PSDB; AAB68049.
DR
XX
PT
     Detecting interaction of cystic fibrosis transmembrane conductance
PT
     regulator (CFTR) polypeptides, useful for screening compounds for
PТ
     treating cystic fibrosis, comprises using yeast dual hybrid assay.
XX
PS
     Disclosure; Page 41-45; 52pp; English.
XX
CC
     The present sequence encodes a human cystic fibrosis transmembrane
CC
     conductance regulator (CFTR) polypeptide. The specification describes a
CC
     method for detecting or determining the interaction of two CFTR
     polypeptides. The method comprises contacting the CFTR polypeptides and
CC
CC
     determining whether the polypeptides interact, where if interaction
CC
     occurs a detectable signal or change is induced in the assay system.
CC
     Polypeptides and polynucleotides that facilitate the interaction of CFTR
CC
     polypeptides are useful for treating cystic fibrosis. Host cells
CC
     comprising the CTFR polynucleotide can be used to model wild-type CFTR
CC
     protein dimerisation, the effect of cystic fibrosis mutations on
CC
     dimerisation and to determine whether a particular mutation of one or
CC
     both the CFTR proteins will effect dimerisation of the CFTR proteins and
CC
     screen for drugs or compounds that can restore or enhance dimerisation of
CC
     CFTR proteins that contain mutations impacting dimerisation
```

```
Sequence 4443 BP; 1363 A; 873 C; 971 G; 1236 T; 0 U; 0 Other;
SQ
                        31.6%; Score 32.2; DB 4; Length 4443;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.75;
                               0; Mismatches
                                                             0; Gaps
                                                                         0;
 Matches 49; Conservative
                                               28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             Db
        1473 TTCTCAGTTTTCCTGGA 1489
RESULT 24
ABX16100
    ABX16100 standard; cDNA; 4443 BP.
XX
AC
    ABX16100;
XX
DT
    08-APR-2003 (first entry)
XX.
    Human cDNA encoding CFTR mutant I539T/R553M/R555K.
DE
XX
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; I539T/R553M/R555K.
ΚW
XX
OS
    Homo sapiens.
     Synthetic.
OS
XX
                    Location/Qualifiers
FH
     Key
                    1. .4443
FT
     CDS
                    /*tag= a
FT
                    /product= "CFTR I539T/R553M/R555K"
FT
                    /transl except= (pos:2496. .2499,aa:Leu)
FT
FT
    mutation
                    replace(1616,T)
                    /*tag= b
FT
                    replace(1656. .1659,CGA)
FT
    mutation
                    /*tag= c
FT
FT
                    replace(1664,G)
    mutation
                    /*tag= d
FT
XX
PN
     US6468793-B1.
XX
PD
     22-OCT-2002.
XX
PF
     22-OCT-1999;
                   99US-00425453.
XX
                   98US-0105444P.
PR
     23-OCT-1998;
XX
PA
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
```

DR

WPI; 2003-182092/18.

```
DR
     P-PSDB; ABG74141.
XX
PT
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Claim 4; Col 79-84; 66pp; English.
XX
     The invention relates to a modified cystic fibrosis transmembrane
CC
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
CC
     a modified CFTR where the modification comprises Ile at position 539
CC
     changed to Thr, Arg at 553 to Met and Arg at 555 Lys
XX
SO
     Sequence 4443 BP; 1364 A; 873 C; 970 G; 1236 T; 0 U; 0 Other;
                          31.6%;
                                  Score 32.2; DB 8; Length 4443;
  Best Local Similarity
                          63.6%;
                                  Pred. No. 0.75;
  Matches
            49; Conservative
                                 0; Mismatches
                                                  28;
                                                       Indels
                                                                 0; Gaps
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                             11 | | | | | | | | | |
                                            Db
         1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
           65 TTGTCACTTTCCGAGGA 81
Qу
              Db
         1473 TTCTCAGTTTTCCTGGA 1489
RESULT 25
ABX16094
ID
     ABX16094 standard; cDNA; 4443 BP.
XX
AC
    ABX16094;
XX
DT
     08-APR-2003 (first entry)
XX
DE
     Human cDNA encoding CFTR mutant I539T.
XX
KW
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; I539T.
XX
OS
     Homo sapiens.
OS
     Synthetic.
XX
FH
     Key
                    Location/Qualifiers
FΤ
                    1. .4443
     CDS
\mathbf{FT}
                    /*tag= a
```

```
FT
                     /product= "CFTR I539T"
                     /transl except= (pos:2496. .2499,aa:Leu)
FT
FT
                     replace (1616, T)
     mutation
FT
                     /*tag= b
XX
     US6468793-B1.
PN
XX
PD
     22-OCT-2002.
XX
PF
     22-OCT-1999;
                   99US-00425453.
XX
PR
     23-OCT-1998;
                   98US-0105444P.
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PI
     Teem JL;
XX
DR
     WPI; 2003-182092/18.
DR
     P-PSDB; ABG74135.
XX
PТ
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Claim 2; Col 11-16; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
CC
     or its biologically active fragment, where expression of the modified
     CFTR protein within a cell results in increased CFTR chloride channel
CC
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
    polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
CC
     a modified CFTR where the modification comprises Ile at position 539
CC
     changed to Thr
XX
SO
     Sequence 4443 BP; 1363 A; 874 C; 971 G; 1235 T; 0 U; 0 Other;
  Query Match
                         31.6%;
                                 Score 32.2; DB 8; Length 4443;
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.75;
 Matches
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                            0;
                                                                0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                             Db
        1413 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
          65 TTGTCACTTTCCGAGGA 81
Qу
              1473 TTCTCAGTTTTCCTGGA 1489
```

RESULT 26 ABX16099

```
ABX16099 standard; cDNA; 4443 BP.
ID
XX
     ABX16099;
AC
XX
DT
     08-APR-2003 (first entry)
XX
DE
     Human cDNA encoding CFTR mutant I539T/G550E.
XX
KW
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
KW
     cystic fibrosis; I539T/G550E.
XX
OS
     Homo sapiens.
OS
     Synthetic.
XX
FH
     Key
                     Location/Qualifiers
FT
                     1. .4443
     CDS
FT
                     /*tag= a
                     /product= "CFTR I539T/G550E"
FT
                     /transl except= (pos:2496. .2499,aa:Leu)
FΤ
FT
                     replace(1616,T)
     mutation
                     /*tag= b
FT
FT
                     replace (1649, A)
     mutation
FT
                     /*tag= c
XX
PN
     US6468793-B1.
XX
PD
     22-OCT-2002.
XX
PF
                    99US-00425453.
     22-OCT-1999;
XX
PR
     23-OCT-1998;
                    98US-0105444P.
XX
PA
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
     WPI; 2003-182092/18.
DR
     P-PSDB; ABG74140.
DR
XX
PT
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
     transmembrane conductance regulator polypeptide.
PT
XX
PS
     Claim 3; Col 69-72; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
```

```
a modified CFTR where the modification comprises Ile at position 539
     changed to Thr and Gly at 550 to Glu
CC
XX
     Sequence 4443 BP; 1364 A; 874 C; 970 G; 1235 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 8; Length 4443;
  Query Match
  Best Local Similarity
                         63.6%; Pred. No. 0.75;
           49; Conservative
                               0; Mismatches
                                                28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
             Db
         1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Qу
          65 TTGTCACTTTCCGAGGA 81
             Db
        1473 TTCTCAGTTTTCCTGGA 1489
RESULT 27
ABX16097
    ABX16097 standard; cDNA; 4443 BP.
ID
XX
AC
    ABX16097;
XX
DT
    08-APR-2003 (first entry)
XX
DΕ
    Human cDNA encoding CFTR mutant R553M.
XX
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; R553M.
XX
os
     Homo sapiens.
os
     Synthetic.
XX
                    Location/Oualifiers
FΗ
     Key
FT
     CDS
                    1. .4443
                    /*tag= a
FT
FT
                    /product= "CFTR R553M"
                    /transl except= (pos:2496. .2499,aa:Leu)
FT
                    replace (1656. .1659, AGA)
FT
    mutation
                    /*tag=b
FT
XX
PN
    US6468793-B1.
XX
PD
    22-OCT-2002.
XX
PF
    22-OCT-1999;
                   99US-00425453.
XX
                   98US-0105444P.
PR
    23-OCT-1998;
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
    Teem JL;
XX
    WPI; 2003-182092/18.
DR
DR
     P-PSDB; ABG74138.
```

```
PT
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Example 2; Col 45-50; 66pp; English.
XX
     The invention relates to a modified cystic fibrosis transmembrane
CC
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
     CFTR protein within a cell results in increased CFTR chloride channel
CC
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
CC
     a modified CFTR where the modification comprises Arg at position 553
CC
     changed to Met
XX
SQ
     Sequence 4443 BP; 1363 A; 872 C; 971 G; 1237 T; 0 U; 0 Other;
  Query Match
                                 Score 32.2; DB 8; Length 4443;
                         31.6%;
                         63.6%;
  Best Local Similarity
                                 Pred. No. 0.75;
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                0; Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
              1473 TTCTCAGTTTTCCTGGA 1489
RESULT 28
ABX16103
    ABX16103 standard; cDNA; 4443 BP.
XX
    ABX16103;
AC
XX
     08-APR-2003 (first entry)
DT
XX
DE
    Human cDNA encoding CFTR mutant I539M/G550E.
XX
KW
    Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; I539T/G550E.
XX
OS
    Homo sapiens.
OS
    Synthetic.
XX
FH
    Key
                    Location/Qualifiers
FT
    CDS
                    1. .4443
FT
                    /*tag= a
FT
                    /product= "CFTR I539T/G550E"
```

```
FT
                    /transl except= (pos:2496. .2499,aa:Leu)
FT
    mutation
                    replace (1617, A)
FT
                    /*tag= b
FT
    mutation
                    replace (1649, A)
FT
                    /*tag= c
XX
    US6468793-B1.
PN
XX
     22-OCT-2002.
PD
XX
PF
     22-OCT-1999;
                   99US-00425453.
XX
PR
     23-OCT-1998;
                   98US-0105444P.
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
    Teem JL;
XX
DR
    WPI; 2003-182092/18.
DR
     P-PSDB; ABG74144.
XX
PT
    Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
     transmembrane conductance regulator polypeptide.
РΤ
XX
PS
     Disclosure; Col 113-118; 66pp; English.
XX
    The invention relates to a modified cystic fibrosis transmembrane
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
CC
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
CC
CC
     a modified CFTR where the modification comprises Ile at position 539
CC
     changed to Met and Gly at 550 to Glu
XX
SQ
     Sequence 4443 BP; 1363 A; 873 C; 971 G; 1236 T; 0 U; 0 Other;
  Query Match
                         31.6%;
                                 Score 32.2; DB 8;
  Best Local Similarity
                                 Pred. No. 0.75;
                         63.6%;
           49; Conservative
                                0; Mismatches
  Matches
                                                 28;
                                                      Indels
                                                                0;
                                                                    Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                             1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              1473 TTCTCAGTTTTCCTGGA 1489
Db
```

```
ID
     ABX16095 standard; cDNA; 4443 BP.
XX
AC
     ABX16095;
XX
DT
     08-APR-2003 (first entry)
XX
DE
     Human cDNA encoding CFTR mutant I539M.
XX
KW
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
KW
     cystic fibrosis; I539M.
XX
OS
     Homo sapiens.
OS
     Synthetic.
XX
FH
                     Location/Qualifiers
     Key
FT
     CDS
                     1. .4443
FT
                     /*tag= a
FT
                     /product= "CFTR I539M"
FT
                     /transl except= (pos:2496. .2499,aa:Leu)
FT
     mutation
                     replace(1617,A)
FT
                     /*tag= b
XX
PN
     US6468793-B1.
XX
PD
     22-OCT-2002.
XX
PF
     22-OCT-1999;
                    99US-00425453.
XX
PR
     23-OCT-1998;
                    98US-0105444P.
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
DR
     WPI; 2003-182092/18.
DR
     P-PSDB; ABG74136.
XX
PT
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Example 1; Col 23-28; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
     a modified CFTR where the modification comprises Ile at position 539
CC
```

```
CC
     changed to Met
XX
SQ
     Sequence 4443 BP; 1362 A; 873 C; 972 G; 1236 T; 0 U; 0 Other;
                         31.6%; Score 32.2; DB 8; Length 4443;
  Query Match
  Best Local Similarity
                         63.6%; Pred. No. 0.75;
  Matches
           49; Conservative 0; Mismatches
                                                 28;
                                                      Indels
                                                                0; Gaps
                                                                            0:
Qу
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
                                            Db
         1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Qу
          65 TTGTCACTTTCCGAGGA 81
              11 | 11 | 11 | 1 | 1 | 1
Db
         1473 TTCTCAGTTTTCCTGGA 1489
RESULT 30
ABX16098
     ABX16098 standard; cDNA; 4443 BP.
XX
AC
     ABX16098;
XX
DT
     08-APR-2003 (first entry)
XX
DE
     Human cDNA encoding CFTR mutant R555K.
XX
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; R555K.
XX
OS
     Homo sapiens.
OS
     Synthetic.
XX
FH
     Key
                    Location/Qualifiers
FT
                    1. .4443
     CDS
FT
                    /*tag= a
FT
                    /product= "CFTR R555K"
FT
                    /transl except= (pos:2496. .2499,aa:Leu)
FT
                    replace (1664, G)
    mutation
FT
                    /*tag= b
XX
PN
    US6468793-B1.
XX
PD
    22-OCT-2002.
XX
PF
    22-OCT-1999;
                   99US-00425453.
XX
PR
    23-OCT-1998;
                   98US-0105444P.
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PI
    Teem JL;
XX
DR
    WPI; 2003-182092/18.
DR
    P-PSDB; ABG74139.
XX
```

```
Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Example 2; Col 57-62; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
CC
     or its biologically active fragment, where expression of the modified
     CFTR protein within a cell results in increased CFTR chloride channel
CC
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
     a modified CFTR where the modification comprises Arg at position 555
CC
CC
     changed to Lys
XX
SQ
     Sequence 4443 BP; 1364 A; 873 C; 970 G; 1236 T; 0 U; 0 Other;
  Query Match
                         31.6%;
                                 Score 32.2; DB 8; Length 4443;
  Best Local Similarity
                         63.6%; Pred. No. 0.75;
  Matches
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                0; Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
        1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 31
ABX16102
ID
    ABX16102 standard; cDNA; 4443 BP.
XX
AC
    ABX16102;
XX
DT
    08-APR-2003 (first entry)
XX
DE
    Human cDNA encoding CFTR mutant I539M/R553M/R555K.
XX
    Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
KW
    cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
    cystic fibrosis; I539M/R553M/R555K.
XX
OS
    Homo sapiens.
OS
    Synthetic.
XX
FΗ
    Key
                    Location/Qualifiers
FT
    CDS
                    1. .4443
FT
                    /*tag= a
FT
                    /product= "CFTR I539M/R553M/R555K"
FT
                    /transl except= (pos:2496. .2499,aa:Leu)
```

PT

```
FT
     mutation
                     replace(1617,A)
FT
                     /*tag= b
                    replace(1656. .1659,CGA)
FT
     mutation
FΤ
                     /*tag= c
FT
     mutation
                    replace (1664, G)
FT
                     /*tag= d
XX
PN
     US6468793-B1.
XX
PD
     22-OCT-2002.
XX
PF
     22-OCT-1999;
                   99US-00425453.
XX
PR
     23-OCT-1998;
                   98US-0105444P.
XX
PΑ
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
     WPI; 2003-182092/18.
DR
DR
     P-PSDB; ABG74143.
XX
PΤ
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
     useful for treating cystic fibrosis, encodes cystic fibrosis
PΤ
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Disclosure; Col 103-106; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
     activity as compared to wild-type CFTR protein. Also included are an
CC
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
CC
     a modified CFTR where the modification comprises Ile at position 539
CC
     changed to Met, Arg at 553 to Met and Arg at 555 Lys
XX
     Sequence 4443 BP; 1363 A; 872 C; 971 G; 1237 T; 0 U; 0 Other;
SQ
  Query Match
                         31.6%;
                                 Score 32.2; DB 8; Length 4443;
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.75;
  Matches
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                           0;
                                                                   Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                             Db
        1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
           65 TTGTCACTTTCCGAGGA 81
Qу
             Db
         1473 TTCTCAGTTTTCCTGGA 1489
```

```
RESULT 32
ABX16096
ID
     ABX16096 standard; cDNA; 4443 BP.
XX
AC
     ABX16096;
XX
DT
     08-APR-2003 (first entry)
XX
DE
     Human cDNA encoding CFTR mutant G550E.
XX
KW
     Human; ss; gene; CFTR; cystic fibrosis; mutant; CFTR chloride channel;
KW
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
     cystic fibrosis; G550E.
XX
OS
     Homo sapiens.
OS.
     Synthetic.
XX
FH
     Key
                     Location/Qualifiers
FT
     CDS
                     1. .4443
FT
                     /*tag= a
                     /product= "CFTR G550E"
FT
FT
                     /transl except= (pos:2496. .2499,aa:Leu)
FT
     mutation
                     replace (1649, A)
FT
                     /*tag=b
XX
PN
     US6468793-B1.
XX
PD
     22-OCT-2002.
XX
PF
     22-OCT-1999;
                    99US-00425453.
XX
PR
     23-OCT-1998;
                    98US-0105444P.
XX
PA
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
     Teem JL;
XX
DR
     WPI; 2003-182092/18.
     P-PSDB; ABG74137.
DR
XX
     Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
PT
     useful for treating cystic fibrosis, encodes cystic fibrosis
PT
     transmembrane conductance regulator polypeptide.
XX
PS
     Example 2; Col 35-38; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
CC
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
CC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
     CFTR polynucleotide is also useful for treating a patient having a
CC
     deficiency or dysfunction in CFTR function. The present sequence encodes
```

```
a modified CFTR where the modification comprises Gly at position 539
CC
     changed to Glu
XX
SQ
     Sequence 4443 BP; 1364 A; 873 C; 970 G; 1236 T; 0 U; 0 Other;
                         31.6%; Score 32.2; DB 8; Length 4443;
  Best Local Similarity 63.6%; Pred. No. 0.75;
  Matches
           49; Conservative
                                0; Mismatches
                                                28; Indels
                                                               0; Gaps
                                                                           0:
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
              1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             11 111 111 1 111
         1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 33
ABX16101
ID
    ABX16101 standard; cDNA; 4443 BP.
XX
AC
    ABX16101;
XX
DT
    08-APR-2003 (first entry)
XX
DΕ
    Human cDNA encoding wild-type CFTR.
XX
KW
    Human; ss; gene; CFTR; cystic fibrosis; CFTR chloride channel;
     cystic fibrosis transmembrane conductance regulator; gene therapy;
KW
KW
    cystic fibrosis.
XX
    Homo sapiens.
OS
XX
FH
                    Location/Qualifiers
    Key
FT
                    1. .4443
    CDS
FT
                    /*tag= a
FT
                    /product= "CFTR"
FT
                    /transl except= (pos:2496. .2499,aa:Leu)
XX
PN
    US6468793-B1.
XX
PD
    22-OCT-2002.
XX
PF
    22-OCT-1999;
                   99US-00425453.
XX
PR
    23-OCT-1998;
                   98US-0105444P.
XX
PA
     (UYFL ) UNIV FLORIDA STATE RES FOUND INC.
XX
PΙ
    Teem JL;
XX
DR
    WPI; 2003-182092/18.
DR
    P-PSDB; ABG74142.
XX
PT
    Novel cystic fibrosis transmembrane conductance regulator polynucleotide
PT
    useful for treating cystic fibrosis, encodes cystic fibrosis
```

CC

```
XX
PS
     Example 1; Col 91-96; 66pp; English.
XX
CC
     The invention relates to a modified cystic fibrosis transmembrane
     conductance regulator (CFTR) polynucleotide encoding a CFTR polypeptide,
СC
     or its biologically active fragment, where expression of the modified
CC
     CFTR protein within a cell results in increased CFTR chloride channel
CC
     activity as compared to wild-type CFTR protein. Also included are an
CC
     isolated cell comprising the CFTR polynucleotide and a polynucleotide
CC
     expression vector comprising the CFTR polynucleotide. The CFTR
CC
     polynucleotide is useful for treating cystic fibrosis by gene therapy and
CC
     for increasing CFTR-mediated chloride channel activity in a cell. The
CC
    CFTR polynucleotide is also useful for treating a patient having a
CC
    deficiency or dysfunction in CFTR function. The present sequence encodes
CC
CC
    wild-type CFTR
XX
     Sequence 4443 BP; 1363 A; 873 C; 971 G; 1236 T; 0 U; 0 Other;
SO
                                 Score 32.2; DB 8; Length 4443;
  Query Match
                         31.6%;
                                 Pred. No. 0.75;
  Best Local Similarity
                         63.6%;
           49; Conservative
                                0; Mismatches
                                                 28; Indels
                                                                0; Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                             1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
          65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
         1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 34
AAZ11643
    AAZ11643 standard; cDNA; 4560 BP.
XX
AC
    AAZ11643;
XX
DT
    26-MAY-2000 (first entry)
XX
DE
    CFTR protein encoding cDNA.
XX
     AAV vector; inverted terminal repeat; ITR; gene therapy; CFTR; TK gene;
KW
     cystic fibrosis transmembrane conductance regulator; cystic fibrosis;
KW
     promoter; HSV; thymidine kinase; chromosome 7q31; ss.
KW
XX
OS
    Homo sapiens.
XX
FH
                     Location/Qualifiers
     Key
FT
     CDS
                     133. .4560
FT
                     /*tag= a
FT
                     /transl except= (pos: 3580. .3582, aa: Ile)
                     /note= "the stop codon is not indicated"
FT
XX
PN
    WO9943789-A1.
XX
PD
     02-SEP-1999.
```

transmembrane conductance regulator polypeptide.

PT

```
XX
PF
     25-FEB-1999;
                   99WO-US004212.
XX
                   98US-0075980P.
     25-FEB-1998;
PR
XX
     (REGC ) UNIV CALIFORNIA.
PA
XX
     Dong J, Kan YW;
PI
XX
     WPI; 1999-550866/46.
DR
     P-PSDB; AAY33968.
DR
XX
     Efficient AAV vectors useful in gene therapy protocols for the treatment
PT
PΤ
     of cystic fibrosis.
XX
     Example 1; Page 33; 34pp; English.
PS
XX
     The invention provides efficient AAV vectors with improved capacity for
CC
     DNA due to the removal of all nucleic acid sequences that are not
CC
     essential for replication (to leave just 2 inverted terminal repeat
CC
CC
     sequences (ITRs)). The AAV vectors may be used for the delivery of
     therapeutic nucleic acids in gene therapy protocols. In particular, they
CC
CC
     may be used to deliver cystic fibrosis (CF) transmembrane conductance
     regulator (CFTR) polynucleotides to the respiratory tract of CF patients
CC
     to rectify mutations in the patients own CFTR genes and restore normal
CC
     function to the chloride channel the gene encodes. The AAV vector lacks
CC
     all nucleic acids that are not essential for replication, therefore
CC
     giving it a greater capacity for exogenous DNA and hence improving the
CC
     efficiency with which it transfects cells. The AAV vectors of the
CC
     invention can efficiently and persistently transfer CFTR polynucleotides
CC
     to the airway epithelium of CF patients without any adverse side effects.
CC
     The present sequence represents a cDNA encoding the CFTR protein
CC
XX
     Sequence 4560 BP; 1397 A; 910 C; 1003 G; 1250 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 2; Length 4560;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.76;
                                0; Mismatches
                                                 28; Indels 0; Gaps
  Matches
            49; Conservative
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                                            1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 11 11 1 111
         1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 35
AAS81827
ID
     AAS81827 standard; cDNA; 4845 BP.
XX
AC
     AAS81827;
XX
DT
     13-FEB-2002 (first entry)
XX
     DNA encoding novel human diagnostic protein #17631.
DE
```

```
XX
     Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW
     food supplement; medical imaging; diagnostic; genetic disorder; ss.
KW
XX
OS
     Homo sapiens.
XX
     WO200175067-A2.
PN
XX
PD
     11-OCT-2001.
XX
     30-MAR-2001; 2001WO-US008631.
PF
XX
     31-MAR-2000; 2000US-00540217.
PR
     23-AUG-2000; 2000US-00649167.
PR
XX
     (HYSE-) HYSEQ INC.
PΑ
XX
                          Tang YT;
PI
     Drmanac RT, Liu C,
XX
     WPI; 2001-639362/73.
DR
DR
     P-PSDB; ABG17640.
XX
     New isolated polynucleotide and encoded polypeptides, useful in
PT
PT
     diagnostics, forensics, gene mapping, identification of mutations
     responsible for genetic disorders or other traits and to assess
PT
PT
     biodiversity.
XX
PS
     Claim 1; SEQ ID NO 17631; 103pp; English.
XX
     The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC
     sequences. (I) is useful as hybridisation probes, polymerase chain
CC
     reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
CC
     and in recombinant production of (II). The polynucleotides are also used
CC
     in diagnostics as expressed sequence tags for identifying expressed
CC
CC
     genes. (I) is useful in gene therapy techniques to restore normal
     activity of (II) or to treat disease states involving (II). (II) is
CC
     useful for generating antibodies against it, detecting or quantitating a
CC
     polypeptide in tissue, as molecular weight markers and as a food
CC
     supplement. (II) and its binding partners are useful in medical imaging
CC
     of sites expressing (II). (I) and (II) are useful for treating disorders
CC
     involving aberrant protein expression or biological activity. The
CC
     polypeptide and polynucleotide sequences have applications in
CC
     diagnostics, forensics, gene mapping, identification of mutations
CC
     responsible for genetic disorders or other traits to assess biodiversity
CC
     and to produce other types of data and products dependent on DNA and
CC
     amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
CC
     coding sequences of the invention. Note: The sequence data for this
CC
     patent did not appear in the printed specification, but was obtained in
CC
CC
     electronic format directly from WIPO at
CC
     ftp.wipo.int/pub/published pct sequences
XX
SQ
     Sequence 4845 BP; 1806 A; 1007 C; 921 G; 1111 T; 0 U; 0 Other;
                                  Score 32.2; DB 5; Length 4845;
  Query Match
                          31.6%;
  Best Local Similarity
                          63.6%;
                                  Prèd. No. 0.77;
                                                   28; Indels
                                                                  0;
                                                                      Gaps
                                                                              0;
            49; Conservative
                                 0; Mismatches
  Matches
```

```
5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             3555 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 3614
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             11 111 111 1 111
        3615 TTCTCAGTTTTCCTGGA 3631
Db
RESULT 36
AAQ13605
    AAQ13605 standard; cDNA; 4894 BP.
ID
XX
AC
    AAQ13605;
XX
DT
    25-MAR-2003 (revised)
    21-NOV-1991 (first entry)
DT
XX
    Cystic fibrosis transmembrane conductance regulator gene.
DE
XX
ΚW
    CFTR; ss.
XX
OS
    Homo sapiens.
XX
FΗ
                    Location/Qualifiers
    Key
                    133. .4575
FT
     CDS
                    /*tag= a
FT
XX
    EP446017-A.
PN
XX
PD
    11-SEP-1991.
XX
PF
     05-MAR-1991;
                   91EP-00301819.
XX
PR
     05-MAR-1990;
                   90US-00488307.
PR
     27-SEP-1990;
                   90US-00589295.
PR
     15-NOV-1990;
                   90US-00613592.
XX
PΑ
     (GENZ ) GENZYME CORP.
XX
    Gregory RJ, Cheng SH, Smith A, Paul S, Hehir KM, Marshall J;
PΙ
XX
DR
    WPI; 1991-268856/37.
DR
     P-PSDB: AAR13894.
XX
     DNA encoding cystic fibrosis trans-membrane regulator - for use in
PT
     treating and diagnosing cystic fibrosis.
PT
XX
PS
     Claim 1; Page 26; 50pp; English.
XX
     The DNA sequence codes for cystic fibrosis transmembrane regulator
CC
     (CFTR). It may be used in gene therapy to obtain in vivo prodn. of CFTR
CC
     in cystic fibrosis patients, and also in the prodn. of CFTR for protein
CC
     replacement therapy. CFTR may also be used in the diagnosis of cystic
CC
     fibrosis by monitoring its presence or absence. See also AAQ13606.
CC
CC
     (Updated on 25-MAR-2003 to correct PA field.)
XX
```

```
Sequence 4894 BP; 1495 A; 960 C; 1094 G; 1345 T; 0 U; 0 Other;
SO
                         31.6%; Score 32.2; DB 2; Length 4894;
  Query Match
                         63.6%; Pred. No. 0.77;
  Best Local Similarity
                                                                           0;
                                0; Mismatches
                                                28; Indels
                                                               0; Gaps
 Matches
           49; Conservative
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                                            1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
QУ
              1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 37
AA068002
    AAO68002 standard; DNA; 5635 BP.
XX
    AAQ68002;
AC
XX
DT
     27-AUG-2003
                 (revised)
     25-MAR-2003
DT
                 (revised)
                 (first entry)
    26-OCT-1995
DT
XX
    Ad2/CFTR-1 nucleotide sequence.
DE
XX
     Recombinant adenovirus; Ad2/CFTR-1; adenovirus 2 serotype; Ela; Elb;
KW
     viral replication; gene expression; gene therapy; cystic fibrosis;
ΚW
     cystic fibrosis transmembrane conductance regulator; CFTR; promoter; E3;
KW
     p19; MHC; class 1; viral latency; pulmonary airway; ds.
KW
XX
     Homo sapiens.
OS
OS
     Unidentified.
XX
                    Location/Qualifiers
FΉ
     Key
                    1. .104
\mathbf{FT}
     repeat region
FT
                    /*tag= a
                    /rpt type= INVERTED
FΤ
FΤ
                    /note= "Represents the origin of replication"
                    190. .380
FT
     enhancer
                    /*tag= b
FT
FT
                    /function= "E1A enhancer and viral packaging domain"
FT
                    380. .500
     promoter
                    /*tag= c
FT
                    /note= "E1A promoter region"
FT
     prim_transcript 499. .5635
FT
FT
                    /*tag= d
                    /note= "Hybrid E1A-CFTR-E1B message"
FT
FT
     5'UTR
                    499. .546
FT
                     /*tag= e
FT
     misc feature
                    547. .595
FT
                    /*tag= f
                    /note= "Synthetic linker sequences"
FT
FT
     misc feature
                    593. .5093
FT
                    /*tag= g
FT
                    /note= "Represents nucleotides 123-4622 of the published
```

```
CFTR cDNA sequence"
FT
FT
     CDS
                     603. .5045
                     /*tag= h
FT
                     /product= "CFTR"
FT
FT
     3'UTR
                     5093. .5635
                     /*tag= i
FT
                     /note= "E1B 3' UTR"
FT
FT
                     5099. .5190
     intron
                     /*tag= j
FT
                     /note= "E1B 3' intron"
FT
FT
     prim transcript 5177. .5635
                     /*tag= k
FT
                     /note= "IX protein mRNA"
FT
                     5201. .5623
FT
     CDS
                     /*tag= 1
FT
                     /product= "IX protein (Hexon-associated protein)"
FT
XX
PN
     WO9412649-A2.
XX
PD
     09-JUN-1994.
XX
                    93WO-US011667.
PF
     02-DEC-1993;
XX
                    92US-00985478.
PR
     03-DEC-1992;
                    93US-00130682.
     01-OCT-1993;
PR
                    93US-00136742.
PR
     13-OCT-1993;
XX
     (GENZ ) GENZYME CORP.
PA
XX
     Gregory RJ, Armentano D, Couture LA,
                                              Smith AE;
PΙ
XX
     WPI: 1994-200277/24.
DR
     P-PSDB; AAR79011, AAR79012.
DR
XX
     Adeno: virus-based gene therapy vectors - esp. useful for gene therapy of
PT
PT
     cystic fibrosis.
XX
     Claim 4; Page 67-80; 167pp; English.
PS
XX
     This sequence represents the nucleotide sequence of the recombinant
CC
     adenovirus Ad2/CFTR-1. This virus is derived from the relatively benign
CC
     adenovirus 2 serotype. The Ela and Elb regions of the viral genome, which
CC
     are involved in the early stages of viral replication have been deleted
CC
     which impairs viral gene expression and viral replication. The cystic
CC
     fibrosis transmembrane conductance regulator (CFTR) coding sequence is
CC
     inserted into the genome in place of the Ela/Elb region and transcription
CC
     of the CFTR sequence is driven by the endogenous Ela promoter. This is a
CC
     moderately strong promoter that is functional in a variety of cells. This
CC
     adenovirus retains the E3 viral coding region. As a consequence the
CC
     length of the adenovirus-CFTR DNA is greater than that of wild type
CC
     adenovirus. This renders the DNA more difficult to package and means that
CC
     the growth of the Ad2/CFTR virus is impaired even in permissive cells
CC
     that provide the missing Ela and Elb functions. The E3 region encodes a
CC
     number of proteins, including p19 which is believed to interact with and
CC
     prevent presentation of MHC class 1 proteins. This property prevents
CC
     recognition of the infected cells and thus may allow viral latency. This
CC
     adenovirus may be administered to the pulmonary airways in the gene
CC
```

```
therapy of cystic fibrosis. (Updated on 25-MAR-2003 to correct PN field.)
CC
     (Updated on 27-AUG-2003 to correct OS field.)
CC
XX
    Sequence 5635 BP; 1619 A; 1142 C; 1324 G; 1550 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 2; Length 5635;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.81;
                                                                          0;
                              0; Mismatches
                                                28; Indels
                                                              0; Gaps
          49; Conservative
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             2015 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 2074
Db
          65 TTGTCACTTTCCGAGGA 81
QУ
             11 111 111 1 111
        2075 TTCTCAGTTTTCCTGGA 2091
RESULT 38
AAQ13053
    AAQ13053 standard; cDNA; 6126 BP.
ID
XX
AC
    AAQ13053;
XX
DT
    14-OCT-1991 (first entry)
XX
    CFTR delta I507.
DE
XX
    Deletion; mutant; diagnosis; antibodies; drug therapy;
KW
KW
    ATP-binding domain; ss.
XX
    Homo sapiens.
OS
XX
                    Location/Qualifiers
FH
    Key
                    133. .4569
FT
     CDS
                    /*tag= a
FT
                    /label= CFTR-mutant
FΤ
FΤ
                    185. .186
    misc feature
                    /*tag= b
FT
                    /label= exon junction
FT
                    296. .297
FT
    misc feature
                    /*tag= c
FT
                    /label= exon junction
FT
                    372. .438
FT
    misc feature
FT
                    /*tag= w
FT
                    /label= membrane-spanning segment
                    405. .406
FT
    misc feature
                    /*tag= d
FT
                    /label= exon junction
FT
                    484. .546
FT
    misc_feature
                    /*tag= x
FT
                    /label= membrane-spanning segment
FT
                    621. .622
FT
    misc feature
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XX
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XX
PD
     25-JUL-1991.
XX
PF
                    90CA-02007699.
     12-JAN-1990;
XX
PR
     12-JAN-1990;
                    90CA-02007699.
PR
     01-MAR-1990;
                    90CA-02011253.
PR
     10-JUL-1990;
                    90CA-02020817.
XX
PA
     (HSCR-) HSC RES DEV CORP.
XX
PΙ
     Tsui LC, Rommens JM,
                            Kerem B;
XX
DR
     WPI: 1991-238022/32.
DR
     P-PSDB; AAR13231.
XX
PT
     Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
XX
PS
     Claim 1; Page 121; 178pp; English.
XX
CC
     The deletion of the 3 bp (ATC) at the I506 or I507 position results in
     the loss of an isoleucine residue from the putative CFTR, within the same
CC
CC
     ATP-binding domain where deltaF508 resides, but it is not evident whether
CC
     this deleted amino acid corresponds to the position 506 or 507. Since the
     506 and 507 positions are repeats, it is at present impossible to
CC
```

```
CC
     followed by a poly(dA) tract. The mutant CF gene when expressed in cells
CC
     of the human body, is associated with altered cell function which
CC
     correlates with the genetic disease cystic fibrosis. See also AAQ13053-72
XX
     Sequence 6126 BP; 1884 A; 1182 C; 1329 G; 1731 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 2; Length 6126;
  Query Match
                         63.6%; Pred. No. 0.83;
  Best Local Similarity
                                                                            0;
  Matches
           49; Conservative
                                0; Mismatches
                                                 28; Indels
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
           65 TTGTCACTTTCCGAGGA 81
Qу
              11 | 11 | 11 | 1 | 11
         1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 39
AAX35553
ID
     AAX35553 standard; DNA; 6126 BP.
XX
AC
    AAX35553;
XX
DT
     08-JUL-1999 (first entry)
XX
DE
     DeltaF508 cystic fibrosis transmembrane conductance regulator DNA.
XX
KW
     Flavone; isoflavone; reservatrol; ascorbic acid; ascorbate salt;
KW
     dehydroascorbic acid; chloride transport; epithelial cell;
KW
     cystic fibrosis; chloride ion conductance;
KW
     cystic fibrosis transmembrane conductance regulator; CFTR;
KW
     chronic bronchitis; asthma; intestinal constipation; ss.
XX
OS
    Homo sapiens.
XX
PN
    WO9918953-A1.
XX
    22-APR-1999.
PD
XX
PF
    16-OCT-1998;
                   98WO-US021887.
XX
PR
     16-OCT-1997;
                   97US-00951912.
XX
PA
     (CHIL-) CHILDREN'S HOSPITAL OAKLAND RES INST.
XX
PI
     Fischer HB, Illek B;
XX
DR
    WPI; 1999-277427/23.
DR
     P-PSDB; AAY02279.
XX
PT
    Use of flavones and isoflavones - for stimulating chloride transport in
PΤ
     epithelial cells and treating cystic fibrosis.
XX
PS
    Disclosure; Page 70-73; 97pp; English.
```

determine in which position the 3 bp deletion occurs. Nucleotide 6126 is

CC

```
CC
    The specification describes compounds comprising flavones/isoflavones,
    reservatrol, ascorbic acid, ascorbate salts and/or dehydroascorbic acid
CC
    which can be used for stimulating chloride transport in epithelial cells
CC
    and treating cystic fibrosis. The compounds can be used to increase
CC
    chloride ion conductance in airway epithelial cells or intestine,
CC
    pancreas, gallbladder, sweat duct, salivary gland or mammary epithelial
CC
    cells. The compounds are useful for treating a patient with cystic
CC
CC
    fibrosis, where the patient's cystic fibrosis transmembrane conductance
    regulator (CFTR) protein has a deletion at position 508 or point mutation
CC
    at 551. They may also be used for treating chronic bronchitis, asthma and
CC
    intestinal constipation. The present sequence encodes a human CTFR
CC
CC
    protein with a F508 deletion mutation
XX
    Sequence 6126 BP; 1886 A; 1181 C; 1330 G; 1729 T; 0 U; 0 Other;
SQ
                         31.6%; Score 32.2; DB 2; Length 6126;
 Ouery Match
 Best Local Similarity
                         63.6%; Pred. No. 0.83;
 Matches
           49; Conservative
                                0; Mismatches
                                                28;
                                                     Indels
                                                               0; Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
Dh
RESULT 40
AAS20529
    AAS20529 standard; DNA; 6126 BP.
ΧX
AC
    AAS20529;
XX
DT
    23-APR-2002 (first entry)
XX
DE
    Human delta-F508-CFTR DNA.
XX
KW
    Human; cystic fibrosis transmembrane conductance regulator; CFTR; gene;
KW
    flavone; isoflavone; chloride transport; epithelial tissue; mucus; ds;
    cystic fibrosis; chronic bronchitis; asthma; delta-F508-CFTR.
KW
XX
OS
    Homo sapiens.
XX
FH
    Key
                    Location/Qualifiers
FT
                    133. .4572
    CDS
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                    /*tag= a
FT
                    /product= "Human delta-F508-CFTR protein"
XX
PN
    US6329422-B1.
XX
PD
    11-DEC-2001.
XX
PF
    16-OCT-1998;
                   98US-00174077.
XX
PR
    16-OCT-1997;
                   97US-00951912.
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XX
PA
     (CHIL-) CHILDREN'S HOSPITAL OAKLAND RES INST.
XX
PΙ
     Fischer H, Illek B;
XX
    WPI; 2002-105224/14.
DR
     P-PSDB; AAU74516.
DR
XX
PT
     Pharmaceutical composition for the treatment of cystic fibrosis comprises
PT
     flavones or isoflavones.
XX
PS
    Disclosure; Col 31-38; 50pp; English.
XX
CC
    The invention relates to a pharmaceutical composition comprising one or
CC
    more compounds such as flavones or isoflavones, capable of stimulating
CC
    chloride transport in epithelial tissues, for treatment of cystic
CC
    fibrosis and other diseases associated with excessive accumulation of
CC
    mucus, e.g. chronic bronchitis and asthma. The active compound increases
CC
     expression of a cystic fibrosis transmembrane conductance regulator
     (CFTR) in an epithelial cell and/or acts as a chemical chaperone that
CC
CC
    increases trafficking of a CFTR to a plasma membrane in an epithelial
CC
    cell. This sequence represents DNA encoding the human delta-F508-CFTR
CC
    mutant polypeptide of the invention
XX
    Sequence 6126 BP; 1886 A; 1181 C; 1330 G; 1729 T; 0 U; 0 Other;
SO
                         31.6%;
                                 Score 32.2; DB 6; Length 6126;
  Query Match
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.83;
           49; Conservative
                                0; Mismatches
 Matches
                                                28;
                                                     Indels
                                                               0; Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             Db
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qy
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 41
ADA37386
    ADA37386 standard; DNA; 6126 BP.
XX
AC
    ADA37386;
XX
DT
    20-NOV-2003 (first entry)
XX
DΕ
    DNA encoding human CFTR F508 deletion mutant.
XX
KW
    ds; gene; cystic fibrosis; chloride transport enhancement;
KW
     epithelial cell; airway epithelial cell; intestinal epithelial cell;
KW
    human; cystic fibrosis transmembrane conductance regulator; CFTR; mutant.
XX
OS
    Homo sapiens.
XX
FH
                    Location/Qualifiers
    Key
FT
    CDS
                    133. .4575
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                     /product= "CFTR F508 deletion mutant"
XX
PN
     US2003096762-A1.
XX
     22-MAY-2003.
PD
XX
     17-OCT-2001; 2001US-00982315.
PF
XX
PR
     16-OCT-1998;
                    98US-00174077.
XX
     (CHIL-) CHILDRENS HOSPITAL OAKLAND.
PA
XX .
PΤ
                Illek B;
     Fischer H,
XX
DR
     WPI; 2003-616312/58.
DR
     P-PSDB; ADA37387.
XX
PT
     Treating cystic fibrosis in a mammal, by administering flavones or
PT
     isoflavones which stimulate chloride secretion, or by administering
PT
     compounds such as resveratrol, ascorbic acid, ascorbate salts or
PT
     dehydroascorbic acid.
XX
PS
     Disclosure; Page 17-20; 34pp; English.
XX
CC
     The invention relates to a method of treating cystic fibrosis in a
CC
     mammal. The method is useful for treating cystic fibrosis in a mammal and
CC
     for enhancing chloride transport in epithelial cells, preferably airway
CC
     epithelial cells or intestinal epithelial cells present in mammals or
CC
     epithelial cells present in pancreas, gallbladder, sweat duct, salivary
     gland or mammary epithelial cells. The present sequence represents the
CC
CC
     human cystic fibrosis transmembrane conductance regulator, CFTR, F508
CC
     deletion mutant.
XX
     Sequence 6126 BP; 1886 A; 1181 C; 1330 G; 1729 T; 0 U; 0 Other;
SQ
  Query Match
                                 Score 32.2; DB 8; Length 6126;
                          31.6%;
  Best Local Similarity
                                 Pred. No. 0.83;
                          63.6%;
            49; Conservative
                                 0; Mismatches
                                                 28; Indels
                                                                 0; Gaps
                                                                             0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
QУ
                             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
           65 TTGTCACTTTCCGAGGA 81
Qy
              11 111 111 1 111
         1605 TTCTCAGTTTTCCTGGA 1621
RESULT 42
AAQ11371
     AAQ11371 standard; DNA; 6127 BP.
XX
AC
    AAQ11371;
XX
DT
     25-MAR-2003
                  (revised)
DT
    22-MAY-1991 (first entry)
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XX
DE
     Mutant cystic fibrosis gene.
XX
KW
     Cystis fibrosis; transmembrane conductance regulatory protein; CFTR;
ΚW
     diagnosis; mutant; ss.
XX
OS
     Homo sapiens.
XX
FΗ
                       Location/Qualifiers
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                       /note= "by primer extension analysis"
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                       297. .405
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                       712. .1001
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\mathbf{FT}
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     WO9102796-A.
XX
PD
     07-MAR-1991.
XX
PF
     22-AUG-1989;
                    89US-00396894.
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PR
     22-AUG-1989;
                    89US-00396894.
PR
     24-AUG-1989;
                    89US-00399945.
PR
     31-AUG-1989;
                    89US-00401609.
XX
PA
     (HSCR-) HSC RES DEV CORP.
PΑ
     (UNMI ) UNIV MICHIGAN.
XX
PΙ
     Tsul LC, Riordan JR, Collins FS,
                                          Rommens JM,
                                                       Jannuzzi MC;
PI
     Kerem BS, Drumm ML, Buckwald M;
XX
DR
     WPI; 1991-087280/12.
     P-PSDB; AAR11602.
DR
XX
PT
     Cystic fibrosis gene - used to produce prods. for screening, detection,
PT
     diagnosis, therapy and studying cystic fibrosis.
XX
PS
     Disclosure; Fig 1; 163pp; English.
XX
CC
     The 3 bp CTT deletion at position 1653-1655 of the normal gene results in
     Phe 508 deletion in the amino acid sequence. The CF gene and its gene
CC
     prod., nucleic acid probes and antibodies to the gene prod. can be used
CC
CC
     for screening and detection of CF carriers, CF diagnosis, prenatal CF
CC
     screening and diagnosis, and gene and drug therapy. The prods. can also
CC
     be used to develop improved methods of treatment and to study the
CC
     disease. See AAQ11046 for the normal CF gene and AAQ11047-48 for CF
CC
     probes. (Updated on 25-MAR-2003 to correct PA field.) (Updated on 25-MAR-
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```
CC
    2003 to correct PI field.)
XX
SQ
     Sequence 6127 BP; 1887 A; 1181 C; 1329 G; 1730 T; 0 U; 0 Other;
  Query Match
                          31.6%; Score 32.2; DB 2; Length 6127;
  Best Local Similarity
                         63.6%; Pred. No. 0.83;
  Matches 49; Conservative
                                0; Mismatches
                                                 28; Indels
                                                                0; Gaps
                                                                            0;
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
              11 111 111 1 111
         1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 43
AA013068
     AAQ13068 standard; DNA; 6128 BP.
XX
AC
    AAQ13068;
XX
DT
    14-OCT-1991 (first entry)
XX
DΕ
    CFTR 556 del A.
XX
KW
    Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
XX
OS
    Homo sapiens.
XX
FH
     Key
                    Location/Qualifiers
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    misc feature
                    296. .297
\mathbf{FT}
                    /*tag= c
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\mathbf{FT}
FT
    misc feature
                    372. .438
FT
                    /*tag= w
FT
                    /label= membrane-spanning segment
FT
    misc feature
                    405. .406
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                    /*tag= d
                    /label= exon junction
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                    484. .545
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                     4373. .4374
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XX
     WO9110734-A.
PN
XX
     25-JUL-1991.
PD
XX
     12-JAN-1990;
                    90CA-02007699.
PF
XX
                    90CA-02007699.
PR
     12-JAN-1990;
                    90CA-02011253.
PR
     01-MAR-1990;
PR
     10-JUL-1990;
                    90CA-02020817.
XX
     (HSCR-) HSC RES DEV CORP.
PA
XX
PΙ
     Tsui LC, Rommens JM,
                            Kerem B;
XX
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     WPI; 1991-238022/32.
     P-PSDB; AAR13304.
DR
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     Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
PT
     fibrosis.
XX
     Claim 2; Page 121; 178pp; English.
PS
XX
     556 del A is a frameshift mutation in exon 4. The mutant CF gene when
CC
     expressed in cells of the human body, is associated with altered cell
CC
     function which correlates with the genetic disease cystic fibrosis. See
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     also AAQ13053-72
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     Sequence 6128 BP; 1884 A; 1183 C; 1329 G; 1732 T; 0 U; 0 Other;
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DT
    14-OCT-1991 (first entry)
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    Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
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    12-JAN-1990;
                    90CA-02007699.
                    90CA-02011253.
PR
     01-MAR-1990;
PR
     10-JUL-1990;
                    90CA-02020817.
XX
PA
     (HSCR-) HSC RES DEV CORP.
XX
PΙ
    Tsui LC, Rommens JM,
                            Kerem B;
XX
DR
    WPI; 1991-238022/32.
DR
     P-PSDB; AAR13308.
XX
PT
    Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
XX
PS
    Claim 2; Page 121; 178pp; English.
XX
CC
     3659 del C is a frameshift mutation in exon 19. The 3659 del C mutation
CC
     results in a shortened polypeptide significantly different from the
CC
     single amino acid deletions or alterations. The mutant CF gene when
CC
     expressed in cells of the human body, is associated with altered cell
CC
     function which correlates with the genetic disease cystic fibrosis. See
CC
    also AAQ13053-72
XX
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     Sequence 6128 BP; 1885 A; 1182 C; 1329 G; 1732 T; 0 U; 0 Other;
  Query Match
                          31.6%; Score 32.2; DB 2; Length 6128;
  Best Local Similarity
                          63.6%; Pred. No. 0.83;
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                               0; Mismatches
                                                 28; Indels
                                                                0; Gaps
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Qу
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        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
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Qу
              11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 45
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ID
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AC
    AAQ13056;
XX
    14-OCT-1991
                (first entry)
DT
XX
DE
    CFTR G178R.
XX
KW
    Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
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OS
    Homo sapiens.
XX
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PR
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PR
     01-MAR-1990;
                    90CA-02020817.
PR
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XX
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PA
XX
     Tsui LC, Rommens JM,
PΙ
                            Kerem B;
XX
     WPI; 1991-238022/32.
DR
     P-PSDB; AAR13234.
DR
XX
     Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
PT
     fibrosis.
XX
     Claim 1; Page 121; 178pp; English.
PS
XX
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CC
     position 664. Nucleotide 6129 is followed by a poly (dA) tract. The
     mutant CF gene when expressed in cells of the human body, is associated
CC
     with altered cell function which correlates with the genetic disease
CC
CC
     cystic fibrosis. See also AAQ13053-72
XX
     Sequence 6129 BP; 1886 A; 1183 C; 1328 G; 1732 T; 0 U; 0 Other;
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                                  Score 32.2; DB 2; Length 6129;
                          31.6%;
  Query Match
                                  Pred. No. 0.83;
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  Best Local Similarity
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Db
RESULT 46
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    14-OCT-1991
                  (first entry)
XX
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DE
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     Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
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OS
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PN
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PR
PR
    10-JUL-1990;
                   90CA-02020817.
XX
     (HSCR-) HSC RES DEV CORP.
PA
XX
PΙ
    Tsui LC, Rommens JM,
                          Kerem B;
XX
DR
    WPI; 1991-238022/32.
DR
    P-PSDB; AAR13297.
XX
PT
    Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
    for producing prods. for diagnosis, screening and therapy of cystic
PT
    fibrosis.
XX
PS
    Claim 1; Page 121; 178pp; English.
XX
CC
    In the S549R, the highly conserved Ser of the nucleotide binding domain
CC
    at position 549 is changed to Arq. The codon change is AGT to AGG. The
CC
    mutant CF gene when expressed in cells of the human body, is associated
CC
    with altered cell function which correlates with the genetic disease
CC
    cystic fibrosis. See also AAQ13053-72
XX
SO
    Sequence 6129 BP; 1885 A; 1183 C; 1330 G; 1731 T; 0 U; 0 Other;
 Query Match
                         31.6%;
                                 Score 32.2; DB 2; Length 6129;
 Best Local Similarity
                         63.6%;
                                 Pred. No. 0.83;
 Matches
           49; Conservative
                                0; Mismatches
                                                 28; Indels
                                                                0; Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                                            Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
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Qу
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RESULT 47
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ΙD
XX
AC
     AAQ13071;
XX
DT
     14-OCT-1991 (first entry)
XX
DE
    CFTR 1717 -1G -> A.
XX
KW
     Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
XX
OS
    Homo sapiens.
XX
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                     Location/Qualifiers
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PN
    WO9110734-A.
XX
    25-JUL-1991.
PD
XX
    12-JAN-1990;
                   90CA-02007699.
PF
XX
                   90CA-02007699.
PR
    12-JAN-1990;
                   90CA-02011253.
PR
    01-MAR-1990;
     10-JUL-1990;
                   90CA-02020817.
PR
XX
     (HSCR-) HSC RES DEV CORP.
PA
XX
PΙ
    Tsui LC, Rommens JM,
                           Kerem B;
XX
    WPI; 1991-238022/32.
DR
XX
    Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
XX
    Claim 2; Page 121; 178pp; English.
PS
XX
CC
    In the 1717 -1G -> A mutation a putative plice mutation is found in front
    of exon 11. This mutation is located at the last nucleotide of the intron
CC
    before exon 11, and is predicted to lead to polypeptides which cannot be
CC
CC
    as yet exactly defined. Nucleotide 6129 is followed by a poly (dA) tract.
    The mutant CF gene when expressed in cells of the human body, is
CC
CC
    associated with altered cell function which correlates with the genetic
    disease cystic fibrosis. See also AAQ13053-72
CC
XX
SO
     Sequence 6129 BP; 1886 A; 1183 C; 1328 G; 1732 T; 0 U; 0 Other;
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  Query Match
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  Best Local Similarity
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                                 Pred. No. 0.83;
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           49; Conservative
                                 0; Mismatches
                                                 28; Indels
                                                                0; Gaps
                                                                            0;
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Db
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     14-OCT-1991 (first entry)
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     CFTR G85E.
XX
     Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
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OS
     Homo sapiens.
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FH
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     CDS
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FT
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                     405. .406
FT
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                     /*tag= d
FT
                     /label= exon junction
FT
FT
     misc feature
                     484. .546
FT
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FT
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FT
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FT
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FT
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                     714. .777
FT
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FT
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FT
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FT
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FT
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FT
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```

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FT
XX
     WO9110734-A.
PN
XX
     25-JUL-1991.
PD
XX
     12-JAN-1990;
                    90CA-02007699.
PF
XX
PR
     12-JAN-1990;
                    90CA-02007699.
PR
     01-MAR-1990;
                    90CA-02011253.
PR
     10-JUL-1990;
                    90CA-02020817.
XX
     (HSCR-) HSC RES DEV CORP.
PΑ
XX
PΙ
     Tsui LC, Rommens JM,
                           Kerem B;
XX
     WPI; 1991-238022/32.
DR
     P-PSDB; AAR13232.
DR
XX
     Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
XX
PS
     Claim 1; Page 121; 178pp; English.
XX
CC
     The G85E mutation in exon3 involves a G to A transition at nucleotide
CC
     position 386. The predicted Gly to Glu amino acid change is associated
CC
     with a group IIb haplotype. The mutation destroys a Hinf1 site.
     Nucleotide 6129 is followed by a poly (dA) tract. The mutant CF gene when
CC
CC
     expressed in cells of the human body, is associated with altered cell
CC
     function which correlates with the genetic disease cystic fibrosis. See
CC
     also AAQ13053-72
XX
SQ
     Sequence 6129 BP; 1886 A; 1183 C; 1328 G; 1732 T; 0 U; 0 Other;
  Query Match
                          31.6%; Score 32.2; DB 2; Length 6129;
  Best Local Similarity
                          63.6%;
                                  Pred. No. 0.83;
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            49; Conservative
                                 0; Mismatches
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                                                       Indels
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                                                                     Gaps
                                                                             0;
Qу
            5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
              Db
         1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Qу
           65 TTGTCACTTTCCGAGGA 81
              11 111 111 1 111
Db
         1605 TTCTCAGTTTTCCTGGA 1621
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ID
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XX
     14-OCT-1991 (first entry)
DT
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    CFTR L1077P.
DE
XX
     Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
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OS
XX
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PR
     01-MAR-1990;
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    10-JUL-1990;
                   90CA-02020817.
PR
XX
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PA
XX
ΡI
    Tsui LC, Rommens JM,
                           Kerem B;
XX
    WPI; 1991-238022/32.
DR
     P-PSDB; AAR13302.
DR
XX
    Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
PT
XX
PS
     Claim 1; Page 121; 178pp; English.
XX
     In the L1077P mutation a T to C change is detected at nucleotide position
CC
     3362. The mutant CF gene when expressed in cells of the human body, is
CC
     associated with altered cell function which correlates with the genetic
CC
     disease cystic fibrosis. See also AAQ13053-72
CC
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SQ
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                                 Score 32.2; DB 2; Length 6129;
  Query Match
                          63.6%;
                                 Pred. No. 0.83;
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                                                 28;
                                                      Indels
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           49; Conservative
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Qу
              1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
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         1605 TTCTCAGTTTTCCTGGA 1621
Db
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RESULT 50 AAQ13063

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АC
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XX
DT
     14-OCT-1991 (first entry)
XX
     CFTR Y563N.
DΕ
XX
KW
     Deletion; mutant; diagnosis; antibodies; drug therapy; ss.
XX
OS
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    12-JAN-1990;
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PR
PR
     01-MAR-1990;
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     10-JUL-1990;
                   90CA-02020817.
PR
XX
     (HSCR-) HSC RES DEV CORP.
PΑ
XX
PΙ
    Tsui LC, Rommens JM,
                           Kerem B;
XX
DR
    WPI; 1991-238022/32.
DR
     P-PSDB; AAR13300.
XX
    Mutant cystic fibrosis trans-membrane conductance regulator gene - used
PT
PT
     for producing prods. for diagnosis, screening and therapy of cystic
PT
     fibrosis.
XX
PS
    Claim 1; Page 121; 178pp; English.
XX
CC
    In the Y563N mutation a T to A change is detected at nucleotide position
     1820 in exon 12. The mutant CF gene when expressed in cells of the human
CC
    body, is associated with altered cell function which correlates with the
CC
CC
     genetic disease cystic fibrosis. See also AAQ13053-72
XX
     Sequence 6129 BP; 1886 A; 1183 C; 1329 G; 1731 T; 0 U; 0 Other;
SO
  Query Match
                         31.6%;
                                 Score 32.2; DB 2; Length 6129;
  Best Local Similarity
                         63.6%;
                                 Pred. No. 0.83;
                                0; Mismatches
                                                                           0;
           49; Conservative
                                                28;
                                                    Indels
                                                               0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
             11 1 1111 11
                             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             Db
        1605 TTCTCAGTTTTCCTGGA 1621
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Search completed: April 29, 2004, 15:06:51 Job time: 53.1699 secs

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OM nucleic - nucleic search, using sw model

April 29, 2004, 14:53:14; Search time 9.33373 Seconds Run on:

(without alignments)

6064.561 Million cell updates/sec

US-09-989-981A-9_COPY_3_104 Title:

Perfect score: 102

Sequence: 1 ctqqtaqqtqaqatctctga.....aacaagctgtcctggaggcc 102

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

682709 seqs, 277475446 residues Searched:

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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6: /cgn2_6/ptodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Res	sult No.	Score	% Query Match	Length	DB	ID	Description
	1	32.2	31.6	420	3	US-09-158-863C-64	Sequence 64, Appl
	2	32.2	31.6	551	2	US-08-647-368A-4	Sequence 4, Appli
С	3	32.2	31.6	558	2	US-08-647-368A-3	Sequence 3, Appli
	4	32.2	31.6	795	4	US-09-866-293-9	Sequence 9, Appli
	5	32.2	31.6	2640	1	US-08-216-971-1	Sequence 1, Appli
	6	32.2	31.6	2640	2	US-08-812-979-1	Sequence 1, Appli
	7	32.2	31.6	2908	3	US-08-487-799-1	Sequence 1, Appli
	8	32.2	31.6	4443	4	US-09-425-453A-1	Sequence 1, Appli
	9	32.2	31.6	4443	4	US-09-425-453A-3	Sequence 3, Appli
	10	32.2	31.6	4443	4	US-09-425-453A-5	Sequence 5, Appli
	11	32.2	31.6	4443	4	US-09-425-453A-7	Sequence 7, Appli

	12	32.2	31.6	4443	4	US-09-425-453A-9	Sequence 9, Appli
	13	32.2	31.6	4443	4	US-09-425-453A-11	Sequence 11, Appl
	14	32.2	31.6	4443	4	US-09-425-453A-13	Sequence 13, Appl
	15	32.2	31.6	4443	4	US-09-425-453A-15	Sequence 15, Appl
	16	32.2	31.6	4443	4	US-09-425-453A-17	Sequence 17, Appl
	17	32.2	31.6	4443	4	US-09-425-453A-19	Sequence 19, Appl
	18	32.2	31.6	4560	3	US-09-256-703-1	Sequence 1, Appli
	19	32.2	31.6	5635	1	US-08-136-742A-3	Sequence 3, Appli
	20	32.2	31.6	5635	3	US-09-248-026-3	Sequence 3, Appli
	21	32.2	31.6	5635	5	PCT-US93-11667-3	Sequence 3, Appli
	22	32.2	31.6	6126	2	US-08-951-912-3	Sequence 3, Appli
	23	32.2	31.6	6126	4	US-09-174-077-3	Sequence 3, Appli
	24	32.2	31.6	6129	1	US-07-637-621-1	Sequence 1, Appli
	25	32.2	31.6	6129	1	US-08-136-742A-1	Sequence 1, Appli
	26	32.2	31.6	6129	1	US-08-135-809A-1	Sequence 1, Appli
	27	32.2	31.6	6129	2	US-08-951-912-1	Sequence 1, Appli
	28	32.2	31.6	6129	2	US-08-951-912-5	Sequence 5, Appli
	29	32.2	31.6	6129	2	US-08-691-605-1	Sequence 1, Appli
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	32	32.2	31.6	6129	4	US-09-174-077-1	Sequence 1, Appli
	33	32.2	31.6	6129	4	US-09-174-077-5	Sequence 5, Appli
	34	32.2	31.6	6129	5	PCT-US93-11667-1	Sequence 1, Appli
	35	32.2	31.6	6130	1	US-08-466-886-16	Sequence 16, Appl
	36	32.2	31.6	6130	2	US-08-604-488-1	Sequence 1, Appli
	37	32.2	31.6	6130	2	US-08-469-461-1	Sequence 1, Appli
	38	32.2	31.6	6130	3	US-07-890-609-1	Sequence 1, Appli
	39	32.2	31.6	6130	3	US-08-030-081-1	Sequence 1, Appli
	40	32.2	31.6	6130	3	US-08-469-617-16	Sequence 16, Appl
	41	32.2	31.6	6146	6	5240846-4	Patent No. 5240846
	42	32.2	31.6	8225	3	US-08-793-618-1	Sequence 1, Appli
	43	32.2	31.6	8225	4	US-09-794-431-1	Sequence 1, Appli
С	44	32.2	31.6	9972	3	US-08-836-022A-3	Sequence 3, Appli
С	45	32.2	31.6	9972	3	US-09-427-048A-3	Sequence 3, Appli
	46	32.2	31.6	12143	4	US-09-423-744A-1	Sequence 1, Appli
	47	32.2	31.6	22846	2	US-08-469-461-3	Sequence 3, Appli
	48	32.2	31.6	22846	3	US-07-890-609-3	Sequence 3, Appli
C	49	30.2	29.6	1251	4	US-09-252-991A-1019	Sequence 1019, Ap
	50	30.2	29.6	2847	4	US-09-252-991A-1036	Sequence 1036, Ap

ALIGNMENTS

RESULT 1

US-09-158-863C-64

- ; Sequence 64, Application US/09158863C
- ; Patent No. 6280978
- ; GENERAL INFORMATION:
- ; APPLICANT: Mitchell, Lloyd G.
- ; APPLICANT: Garcia-Blanco, Mariano A.
- ; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
- ; TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
- ; FILE REFERENCE: 31304-B-A
- ; CURRENT APPLICATION NUMBER: US/09/158,863C
- ; CURRENT FILING DATE: 1998-09-23
- ; PRIOR APPLICATION NUMBER: 09/133,717

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PRIOR FILING DATE: 1998-08-13
   PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
  PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
  PRIOR APPLICATION NUMBER: 60/008,317
  PRIOR FILING DATE: 1995-12-07
  NUMBER OF SEO ID NOS: 68
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 64
   LENGTH: 420
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: trans-spliced product comprising cystic fibrosis
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US-09-158-863C-64
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             Db
         188 TTCTCAGTTTTCCTGGA 204
RESULT 2
US-08-647-368A-4
; Sequence 4, Application US/08647368A
; Patent No. 5928906
  GENERAL INFORMATION:
    APPLICANT: Koster, Hubert
    APPLICANT: Van de Boom, Dirk
    APPLICANT: Ruppert, Andreas
    TITLE OF INVENTION: PROCESS FOR DIRECT SEQUENCING DURING
    TITLE OF INVENTION: TEMPLATE AMPLIFICATION
    NUMBER OF SEQUENCES: 4
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: FOLEY, HOAG & ELIOT LLP
      STREET: One Post Office Square
      CITY: Boston
      STATE: MA
      COUNTRY: USA
      ZIP: 02109-2170
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/647,368A
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FILING DATE: 09-MAY-1996
      CLASSIFICATION: 435
    ATTORNEY/AGENT INFORMATION:
      NAME: Arnold, Beth E.
      REGISTRATION NUMBER: 35,430
      REFERENCE/DOCKET NUMBER: SQA-020.01
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 617-832-1000
      TELEFAX: 617-832-7000
  INFORMATION FOR SEQ ID NO: 4:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 551 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: DNA
US-08-647-368A-4
  Query Match
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  Best Local Similarity 63.6%; Pred. No. 0.023;
 Matches 49; Conservative 0; Mismatches 28; Indels
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Db ·
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Qу
             Db
         145 TTCTCAGTTTTCCTGGA 161
RESULT 3
US-08-647-368A-3/c
; Sequence 3, Application US/08647368A
; Patent No. 5928906
  GENERAL INFORMATION:
    APPLICANT: Koster, Hubert
    APPLICANT: Van de Boom, Dirk
    APPLICANT: Ruppert, Andreas
    TITLE OF INVENTION: PROCESS FOR DIRECT SEQUENCING DURING
    TITLE OF INVENTION: TEMPLATE AMPLIFICATION
    NUMBER OF SEQUENCES: 4
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: FOLEY, HOAG & ELIOT LLP
      STREET: One Post Office Square
      CITY: Boston
      STATE: MA
      COUNTRY: USA
      ZIP: 02109-2170
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/647,368A
      FILING DATE: 09-MAY-1996
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ATTORNEY/AGENT INFORMATION:
      NAME: Arnold, Beth E.
      REGISTRATION NUMBER: 35,430
      REFERENCE/DOCKET NUMBER: SQA-020.01
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 617-832-1000
      TELEFAX: 617-832-7000
  INFORMATION FOR SEQ ID NO: 3:
   SEQUENCE CHARACTERISTICS:
      LENGTH: 558 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: DNA
US-08-647-368A-3
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Qу
            432 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 373
        65 TTGTCACTTTCCGAGGA 81
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            372 TTCTCAGTTTTCCTGGA 356
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RESULT 4
US-09-866-293-9
; Sequence 9, Application US/09866293
; Patent No. 6607911
; GENERAL INFORMATION:
 APPLICANT: Gordon, Joan
  APPLICANT: Rundell, Clark
; TITLE OF INVENTION: COMPOSITIONS AND METHODS RELATING TO CONTROL DNA
CONSTRUCT
; FILE REFERENCE: 053689-5010
 CURRENT APPLICATION NUMBER: US/09/866,293
; CURRENT FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 10
 SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 795
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-866-293-9
                      31.6%; Score 32.2; DB 4; Length 795;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.026;
 Matches 49; Conservative 0; Mismatches 28; Indels
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CLASSIFICATION: 435

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RESULT 5
US-08-216-971-1
; Sequence 1, Application US/08216971
; Patent No. 5639661
  GENERAL INFORMATION:
    APPLICANT: Welsh, Michael J.
    APPLICANT: Sheppard, David N.
    TITLE OF INVENTION: NOVEL GENES AND PROTEINS FOR TREATING
    TITLE OF INVENTION: CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 2
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: LAHIVE & COCKFIELD
      STREET: 60 State Street, suite #510
      CITY: Boston
      STATE: Massachusetts
      COUNTRY: USA
      ZIP: 02109-1875
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/216,971
      FILING DATE: 23-MAR-1994
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
      NAME: Arnold, Beth E.
      REGISTRATION NUMBER: 35,430
      REFERENCE/DOCKET NUMBER: UIZ-011
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 227-7400
      TELEFAX: (617) 227-5941
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 2640 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..2640
US-08-216-971-1
                        31.6%; Score 32.2; DB 1; Length 2640;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.042;
  Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
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Db
RESULT 6
US-08-812-979-1
; Sequence 1, Application US/08812979
; Patent No. 5958893
; GENERAL INFORMATION:
    APPLICANT: Welsh, Michael J.
    APPLICANT: Sheppard, David N.
    TITLE OF INVENTION: NOVEL GENES AND PROTEINS FOR TREATING
    TITLE OF INVENTION: CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 2
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: LAHIVE & COCKFIELD
      STREET: 60 State Street, suite #510
      CITY: Boston
      STATE: Massachusetts
      COUNTRY: USA
      ZIP: 02109-1875
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/08/812,979
      FILING DATE:
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 08/216,971
      FILING DATE:
    ATTORNEY/AGENT INFORMATION:
      NAME: Arnold, Beth E.
      REGISTRATION NUMBER: 35,430
      REFERENCE/DOCKET NUMBER: UIZ-011
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 227-7400
      TELEFAX: (617) 227-5941
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INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2640 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA

FEATURE:

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Matches 49; Conservative 0; Mismatches 28; Indels
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US-08-487-799-1
; Sequence 1, Application US/08487799C
; Patent No. 6010908
; GENERAL INFORMATION:
; APPLICANT: Gruenert, Deiter C.
  APPLICANT: Kunzelmann, Karl
 TITLE OF INVENTION: GENE THERAPY BY SMALL FRAGMENTS HOMOLOGOUS REPLACEMENT
; FILE REFERENCE: 480.18-1(HV)
  CURRENT APPLICATION NUMBER: US/08/487,799C
  CURRENT FILING DATE: 1995-06-07
  EARLIER APPLICATION NUMBER: 07/933,471
 EARLIER FILING DATE: 1992-08-21
  EARLIER APPLICATION NUMBER: 08/409,544
  EARLIER FILING DATE: 1995-03-24
; NUMBER OF SEQ ID NOS: 87
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 2908
   TYPE: DNA
   ORGANISM: human
US-08-487-799-1
  Query Match 31.6%; Score 32.2; DB 3; Length 2908; Best Local Similarity 63.6%; Pred. No. 0.044;
 Matches 49; Conservative 0; Mismatches 28; Indels
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RESULT 8
US-09-425-453A-1
; Sequence 1, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
 TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
; CURRENT APPLICATION NUMBER: US/09/425,453A
; CURRENT FILING DATE: 1999-10-22
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PRIOR APPLICATION NUMBER: 60/105,444
  PRIOR FILING DATE: 1998-10-23
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: gene
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US-09-425-453A-1
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 Best Local Similarity 63.6%; Pred. No. 0.052;
 Matches 49; Conservative
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Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             11 111 111 1 111
        1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 9
US-09-425-453A-3
; Sequence 3, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
 APPLICANT: Teem. John L.
  TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
  FILE REFERENCE: FSU-99XC1
  CURRENT APPLICATION NUMBER: US/09/425,453A
  CURRENT FILING DATE: 1999-10-22
  PRIOR APPLICATION NUMBER: 60/105,444
  PRIOR FILING DATE: 1998-10-23
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: gene
   LOCATION: (1)..(4443)
US-09-425-453A-3
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  Best Local Similarity 63.6%; Pred. No. 0.052;
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RESULT 10
US-09-425-453A-5
; Sequence 5, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
 TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
; CURRENT APPLICATION NUMBER: US/09/425,453A
  CURRENT FILING DATE: 1999-10-22
; PRIOR APPLICATION NUMBER: 60/105,444
; PRIOR FILING DATE: 1998-10-23
; NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-425-453A-5
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Db
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Qу
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Db
RESULT 11
US-09-425-453A-7
; Sequence 7, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
  TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
; CURRENT APPLICATION NUMBER: US/09/425,453A
; CURRENT FILING DATE: 1999-10-22
; PRIOR APPLICATION NUMBER: 60/105,444
; PRIOR FILING DATE: 1998-10-23
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 7
   LENGTH: 4443
   TYPE: DNA
  ORGANISM: Homo sapiens
  FEATURE:
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NAME/KEY: gene
  LOCATION: (1)..(4443)
US-09-425-453A-7
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 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.052;
 Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
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QУ
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RESULT 12
US-09-425-453A-9
; Sequence 9, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
 TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
  FILE REFERENCE: FSU-99XC1
  CURRENT APPLICATION NUMBER: US/09/425,453A
; CURRENT FILING DATE: 1999-10-22
; PRIOR APPLICATION NUMBER: 60/105,444
 PRIOR FILING DATE: 1998-10-23
; NUMBER OF SEQ ID NOS: 20
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   ORGANISM: Homo sapiens
   FEATURE:
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   NAME/KEY: gene
   LOCATION: (1)..(4443)
US-09-425-453A-9
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  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.052;
  Matches 49; Conservative 0; Mismatches 28; Indels
                                                         0; Gaps
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RESULT 13
US-09-425-453A-11
; Sequence 11, Application US/09425453A
; Patent No. 6468793
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; GENERAL INFORMATION:
 APPLICANT: Teem, John L.
 TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
 CURRENT APPLICATION NUMBER: US/09/425,453A
  CURRENT FILING DATE: 1999-10-22
 PRIOR APPLICATION NUMBER: 60/105,444
  PRIOR FILING DATE: 1998-10-23
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 11
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-425-453A-11
                       31.6%; Score 32.2; DB 4; Length 4443;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.052;
                                                          0; Gaps
 Matches 49; Conservative 0; Mismatches 28; Indels
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
            1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
         65 TTGTCACTTTCCGAGGA 81
Qy
            1473 TTCTCAGTTTTCCTGGA 1489
RESULT 14
US-09-425-453A-13
; Sequence 13, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
  APPLICANT: Teem, John L.
  TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
; CURRENT APPLICATION NUMBER: US/09/425,453A
; CURRENT FILING DATE: 1999-10-22
; PRIOR APPLICATION NUMBER: 60/105,444
; PRIOR FILING DATE: 1998-10-23
; NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-425-453A-13
                       31.6%; Score 32.2; DB 4; Length 4443;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.052;
                                                          0; Gaps 0;
 Matches 49; Conservative 0; Mismatches 28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
QУ
            1413 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
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65 TTGTCACTTTCCGAGGA 81
Qу
             11 111 111 1 111
        1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 15
US-09-425-453A-15
; Sequence 15, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
; TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
; CURRENT APPLICATION NUMBER: US/09/425,453A
; CURRENT FILING DATE: 1999-10-22
  PRIOR APPLICATION NUMBER: 60/105,444
  PRIOR FILING DATE: 1998-10-23
  NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 15
   LENGTH: 4443
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
; NAME/KEY: gene
   LOCATION: (1)..(4443)
US-09-425-453A-15
                        31.6%; Score 32.2; DB 4; Length 4443;
  Query Match
                        63.6%; Pred. No. 0.052;
  Best Local Similarity
  Matches 49; Conservative 0; Mismatches 28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                          11 1 11 11
        1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
        65 TTGTCACTTTCCGAGGA 81
Qу
             1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 16
US-09-425-453A-17
; Sequence 17, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
  TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
 FILE REFERENCE: FSU-99XC1
  CURRENT APPLICATION NUMBER: US/09/425,453A
  CURRENT FILING DATE: 1999-10-22
  PRIOR APPLICATION NUMBER: 60/105,444
 PRIOR FILING DATE: 1998-10-23
 NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 17
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LENGTH: 4443

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TYPE: DNA
  ORGANISM: Homo sapiens
US-09-425-453A-17
              31.6%; Score 32.2; DB 4; Length 4443;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.052;
 Matches 49: Conservative 0: Mismatches 28: Indels
                                                                    0;
                                                          0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
QУ
            1413 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
          65 TTGTCACTTTCCGAGGA 81
QУ
            11 111 111 1 111
        1473 TTCTCAGTTTTCCTGGA 1489
RESULT 17
US-09-425-453A-19
; Sequence 19, Application US/09425453A
; Patent No. 6468793
; GENERAL INFORMATION:
; APPLICANT: Teem, John L.
  TITLE OF INVENTION: CFTR Genes and Proteins for Cystic Fibrosis Gene Therapy
; FILE REFERENCE: FSU-99XC1
  CURRENT APPLICATION NUMBER: US/09/425,453A
  CURRENT FILING DATE: 1999-10-22
; PRIOR APPLICATION NUMBER: 60/105,444
; PRIOR FILING DATE: 1998-10-23
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 19
  LENGTH: 4443
   TYPE: DNA
  ORGANISM: Homo sapiens
US-09-425-453A-19
                       31.6%; Score 32.2; DB 4; Length 4443;
 Query Match
 Query Match 31.6%; Score 32.2; DB 4
Best Local Similarity 63.6%; Pred. No. 0.052;
 Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps
                                                                    0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1413 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1472
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
            1473 TTCTCAGTTTTCCTGGA 1489
Db
RESULT 18
US-09-256-703-1
; Sequence 1, Application US/09256703
; Patent No. 6294379
; GENERAL INFORMATION:
; APPLICANT: Dong, Jian-yun
; APPLICANT: Kan, Yuet Wai
```

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APPLICANT: The Regents of the University of California
  TITLE OF INVENTION: Efficient AAV Vectors
  FILE REFERENCE: 023070-084910US
  CURRENT APPLICATION NUMBER: US/09/256,703
  CURRENT FILING DATE: 1999-02-24
  PRIOR APPLICATION NUMBER: US 60/075,980
  PRIOR FILING DATE: 1998-02-25
  NUMBER OF SEQ ID NOS: 7
  SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
   LENGTH: 4560
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: truncated cystic fibrosis transmembrane
   OTHER INFORMATION: conductance regulator (CFTR) polynucleotide
   OTHER INFORMATION: encoding a functional CFTR polypeptide
   NAME/KEY: CDS
   LOCATION: (133)..(4560)
US-09-256-703-1
                        31.6%; Score 32.2; DB 3; Length 4560;
 Query Match
                        63.6%; Pred. No. 0.052;
 Best Local Similarity
                               0; Mismatches
                                               28; Indels
                                                             0; Gaps
                                                                         0;
          49; Conservative
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                           1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 19
US-08-136-742A-3
; Sequence 3, Application US/08136742A
; Patent No. 5670488
  GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
    APPLICANT: A.E.
    TITLE OF INVENTION: GENE THERAPY FOR CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 10
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: BRUMBAUGH, GRAVES, DONOHUE & RAYMOND
      STREET: 30 ROCKEFELLER PLAZA
      CITY: NEW YORK
      STATE: NEW YORK
      COUNTRY: USA
      ZIP: 10112
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/136,742A
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FILING DATE: 02-DEC-1993
      CLASSIFICATION: 514
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 07/985,478
      FILING DATE: 02-DEC-1992
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
     NAME: Seide, Rochelle K.
      REGISTRATION NUMBER: 32,300
      REFERENCE/DOCKET NUMBER: A30668 (Genzyme Dkt. IG4-9.11)
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (212) 408-2500
      TELEFAX: (212) 765-2519
  INFORMATION FOR SEQ ID NO: 3:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 5635 base pairs
      TYPE: nucleic acid
     STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
US-08-136-742A-3
                        31.6%; Score 32.2; DB 1; Length 5635;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.057;
          49; Conservative 0; Mismatches 28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             2015 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 2074
Db
          65 TTGTCACTTTCCGAGGA 81
Qy
             2075 TTCTCAGTTTTCCTGGA 2091
Db
RESULT 20
US-09-248-026-3
; Sequence 3, Application US/09248026
; Patent No. 6093567
  GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
    APPLICANT: A.E.
    TITLE OF INVENTION: ADENOVIRUS VECTORS FOR GENE THERAPY
    NUMBER OF SEQUENCES: 10
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: BAKER & BOTTS, L.L.P.
      STREET: 30 ROCKEFELLER PLAZA
      CITY: NEW YORK
      STATE: NEW YORK
      COUNTRY: USA
      ZIP: 10112
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
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APPLICATION NUMBER: US/09/248,026
      FILING DATE: 10-FEB-1999
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 08/895,194
      FILING DATE: 16-JUL-1997
    ATTORNEY/AGENT INFORMATION:
      NAME: Seide, Rochelle K.
      REGISTRATION NUMBER: 32,300
      REFERENCE/DOCKET NUMBER: A30668-C
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (212) 705-5000
      TELEFAX: (212) 705-5020
  INFORMATION FOR SEQ ID NO: 3:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 5635 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
US-09-248-026-3
                        31.6%; Score 32.2; DB 3; Length 5635;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.057;
 Matches 49; Conservative 0; Mismatches 28; Indels
                                                            0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             2015 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 2074
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             2075 TTCTCAGTTTTCCTGGA 2091
RESULT 21
PCT-US93-11667-3
; Sequence 3, Application PC/TUS9311667
  GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
    APPLICANT: A.E.
    TITLE OF INVENTION: GENE THERAPY FOR CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 9
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: LAHIVE & COCKFIELD
      STREET: 60 STATE STREET, SUITE 510
      CITY: BOSTON
      STATE: MASSACHUSETTS
      COUNTRY: USA
      ZIP: 02109
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: PCT/US93/11667
```

```
FILING DATE: 02-DEC-1993
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 07/985,478
      FILING DATE: 02-DEC-1992
    ATTORNEY/AGENT INFORMATION:
      NAME: Hanley, Elizabeth A.
      REGISTRATION NUMBER: 33,505
      REFERENCE/DOCKET NUMBER: NZI-014CP2
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 227-7400
      TELEFAX: (617) 227-5941
  INFORMATION FOR SEQ ID NO: 3:
    SEQUENCE CHARACTERISTICS:
    LENGTH: 5635 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
PCT-US93-11667-3
                        31.6%; Score 32.2; DB 5; Length 5635;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.057;
 Matches 49; Conservative 0; Mismatches
                                              28; Indels
                                                            0; Gaps
                                                                        0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
QУ
             2015 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 2074
          65 TTGTCACTTTCCGAGGA 81
Qу
            2075 TTCTCAGTTTTCCTGGA 2091
RESULT 22
US-08-951-912-3
; Sequence 3, Application US/08951912
; Patent No. 5972995
  GENERAL INFORMATION:
    APPLICANT: Fischer, Horst
    APPLICANT: Illek, Beate
    TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC
    TITLE OF INVENTION: FIBROSIS THERAPY
    NUMBER OF SEQUENCES: 6
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: SEED and BERRY LLP
      STREET: 6300 Columbia Center, 701 Fifth Avenue
      CITY: Seattle
      STATE: Washington
      COUNTRY: USA
      ZIP: 98104
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
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APPLICATION NUMBER: US/08/951,912
      FILING DATE: 16-OCT-1997
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
      NAME: Maki, David J.
      REGISTRATION NUMBER: 31,392
      REFERENCE/DOCKET NUMBER: 200116.403
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (206) 622-4900
      TELEFAX: (206) 682-6031
  INFORMATION FOR SEQ ID NO: 3:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6126 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4569
US-08-951-912-3
                       31.6%; Score 32.2; DB 2; Length 6126;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps
                                                                       0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qγ
            1605 TTCTCAGTTTTCCTGGA 1621
RESULT 23
US-09-174-077-3
; Sequence 3, Application US/09174077
; Patent No. 6329422
; GENERAL INFORMATION:
; APPLICANT: Fischer, Horst
 APPLICANT: Illek, Beate
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
 FILE REFERENCE: 200116.403C1
  CURRENT APPLICATION NUMBER: US/09/174,077
; CURRENT FILING DATE: 1998-10-16
; EARLIER APPLICATION NUMBER: US 08/951,912
; EARLIER FILING DATE: 1997-10-16
; NUMBER OF SEQ ID NOS: 6
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
   LENGTH: 6126
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-174-077-3
                       31.6%; Score 32.2; DB 4; Length 6126;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
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0; Gaps
 Matches
           49; Conservative 0; Mismatches 28; Indels
                                                                       0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qy
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 24
US-07-637-621-1
; Sequence 1, Application US/07637621
; Patent No. 5407796
  GENERAL INFORMATION:
    APPLICANT: cutting, gary
    APPLICANT: antonarakis, stylianos e
    APPLICANT: kazazian jr., haig h
    TITLE OF INVENTION: CYSTIC FIBROSIS MUTATION CLUSTER
    NUMBER OF SEQUENCES: 4
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Banner, Birch, McKie and Beckett
      STREET: 1001 G Street, N.W.
      CITY: Washington, D.C.
      COUNTRY: USA
      ZIP: 20001
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
;
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/637,621
     FILING DATE: 19910104
      CLASSIFICATION: 435
    ATTORNEY/AGENT INFORMATION:
;
      NAME: kagan, sarah a
      REGISTRATION NUMBER: 32,141
      REFERENCE/DOCKET NUMBER: 1107.030010
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 202-508-9100
      TELEFAX: 202-508-9100
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
      TYPE: NUCLEIC ACID
      STRANDEDNESS: double
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    HYPOTHETICAL: NO
    ANTI-SENSE: NO
    ORIGINAL SOURCE:
      ORGANISM: Homo sapiens
US-07-637-621-1
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31.6%; Score 32.2; DB 1; Length 6129;

Query Match

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Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
            Db
       1605 TTCTCAGTTTTCCTGGA 1621
RESULT 25
US-08-136-742A-1
; Sequence 1, Application US/08136742A
; Patent No. 5670488
; GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
    APPLICANT: A.E.
    TITLE OF INVENTION: GENE THERAPY FOR CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 10
    CORRESPONDENCE ADDRESS:
     ADDRESSEE: BRUMBAUGH, GRAVES, DONOHUE & RAYMOND
     STREET: 30 ROCKEFELLER PLAZA
     CITY: NEW YORK
     STATE: NEW YORK
    COUNTRY: USA
     ZIP: 10112
    COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
     OPERATING SYSTEM: PC-DOS/MS-DOS
     SOFTWARE: ASCII
   CURRENT APPLICATION DATA:
    APPLICATION NUMBER: US/08/136,742A
     FILING DATE: 02-DEC-1993
     CLASSIFICATION: 514
    PRIOR APPLICATION DATA:
    APPLICATION NUMBER: US 07/985,478
;
     FILING DATE: 02-DEC-1992
;
     CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
    NAME: Seide, Rochelle K.
     REGISTRATION NUMBER: 32,300
     REFERENCE/DOCKET NUMBER: A30668 (Genzyme Dkt. IG4-9.11)
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (212) 408-2500
      TELEFAX: (212) 765-2519
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
     LENGTH: 6129 base pairs
     TYPE: nucleic acid
     STRANDEDNESS: single
     TOPOLOGY: linear
   MOLECULE TYPE: cDNA
   FEATURE:
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NAME/KEY: CDS

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LOCATION: 133..4572
US-08-136-742A-1
 Query Match 31.6%; Score 32.2; DB 1; Length 6129; Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches 49; Conservative 0; Mismatches 28; Indels 0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
RESULT 26
US-08-135-809A-1
; Sequence 1, Application US/08135809A
; Patent No. 5688677
; GENERAL INFORMATION:
    APPLICANT: CHENG, SENG H.
    APPLICANT: DITULLIO, PAUL
    APPLICANT: EBERT, KARL M.
    APPLICANT: MEADE, HARRY M.
    APPLICANT: SMITH, ALAN E.
    TITLE OF INVENTION: DEOXYRIBONUCLEIC ACIDS CONTAINING
    TITLE OF INVENTION: INACTIVATED HORMONE RESPONSIVE ELEMENTS
    NUMBER OF SEQUENCES: 9
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: GENZYME CORPORATION
      STREET: ONE MOUNTAIN ROAD
      CITY: FRAMINGHAM
      STATE: MASSACHUSETTS
      COUNTRY: USA
      ZIP: 01701
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
    APPLICATION NUMBER: US/08/135,809A
      FILING DATE: 13-OCT-1993
     CLASSIFICATION: 800
    ATTORNEY/AGENT INFORMATION:
     NAME: LASSEN, ELIZABETH
      REGISTRATION NUMBER: 31,845
      REFERENCE/DOCKET NUMBER: IG4-9.12
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (508) 872-8400
      TELEFAX: (508) 872-5415
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
;
      LENGTH: 6129 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
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TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
US-08-135-809A-1
 Query Match
                        31.6%; Score 32.2; DB 1; Length 6129;
 Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches
         49; Conservative 0; Mismatches 28; Indels 0; Gaps
                                                                        0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qv
             Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qγ
             1605 TTCTCAGTTTTCCTGGA 1621
RESULT 27
US-08-951-912-1
; Sequence 1, Application US/08951912
; Patent No. 5972995
  GENERAL INFORMATION:
    APPLICANT: Fischer, Horst
    APPLICANT: Illek, Beate
    TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC
    TITLE OF INVENTION: FIBROSIS THERAPY
    NUMBER OF SEQUENCES:
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: SEED and BERRY LLP
      STREET: 6300 Columbia Center, 701 Fifth Avenue
      CITY: Seattle
      STATE: Washington
      COUNTRY: USA
      ZIP: 98104
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/951,912
      FILING DATE: 16-OCT-1997
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
;
      NAME: Maki, David J.
      REGISTRATION NUMBER: 31,392
      REFERENCE/DOCKET NUMBER: 200116.403
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (206) 622-4900
      TELEFAX: (206) 682-6031
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
      TYPE: nucleic acid
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STRANDEDNESS: single
      TOPOLOGY: linear
    FEATURE:
      NAME/KEY:
                CDS
      LOCATION: 133..4572
US-08-951-912-1
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 Matches
         49; Conservative
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Qу
             Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 28
US-08-951-912-5
; Sequence 5, Application US/08951912
; Patent No. 5972995
  GENERAL INFORMATION:
    APPLICANT: Fischer, Horst
    APPLICANT: Illek, Beate
    TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC
    TITLE OF INVENTION: FIBROSIS THERAPY
    NUMBER OF SEQUENCES: 6
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: SEED and BERRY LLP
      STREET: 6300 Columbia Center, 701 Fifth Avenue
      CITY: Seattle
      STATE: Washington
      COUNTRY: USA
      ZIP: 98104
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/951,912
      FILING DATE: 16-OCT-1997
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
      NAME: Maki, David J.
      REGISTRATION NUMBER: 31,392
      REFERENCE/DOCKET NUMBER: 200116.403
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (206) 622-4900
      TELEFAX: (206) 682-6031
  INFORMATION FOR SEQ ID NO: 5:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
      TYPE: nucleic acid
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STRANDEDNESS: single
      TOPOLOGY: linear
US-08-951-912-5
  Query Match
                       31.6%; Score 32.2; DB 2; Length 6129;
  Best Local Similarity 63.6%; Pred. No. 0.059;
  Matches 49; Conservative 0; Mismatches 28; Indels
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                                                                        0;
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Qу
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 29
US-08-691-605-1
; Sequence 1, Application US/08691605
; Patent No. 5981714
  GENERAL INFORMATION:
    APPLICANT: Cheng, Seng H., Marshall, John, Gregory, Richard J.
    APPLICANT: and Rafter, Patrick. W.
    TITLE OF INVENTION: ANTIBODIES SPECIFIC FOR CYSTIC FIBROSIS
    TITLE OF INVENTION: TRANSMEMBRANE CONDUCTANCE REGULATOR AND USES
    TITLE OF INVENTION: THEREFOR
    NUMBER OF SEQUENCES: 2
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: LAHIVE & COCKFIELD
      STREET: 60 STATE STREET, SUITE 510
      CITY: BOSTON
      STATE: MASSACHUSETTS
      COUNTRY: USA
      ZIP: 02109
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/691,605
      FILING DATE:
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 08/114,950
      FILING DATE:
      CLASSIFICATION:
    ATTORNEY/AGENT INFORMATION:
      NAME: Hanley, Elizabeth A.
      REGISTRATION NUMBER: 33,505
      REFERENCE/DOCKET NUMBER: NZI-029
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 227-7400
      TELEFAX: (617) 227-5941
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
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LENGTH: 6129 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
US-08-691-605-1
  Query Match 31.6%; Score 32.2; DB 2; Length 6129; Best Local Similarity 63.6%; Pred. No. 0.059;
  Matches 49; Conservative 0; Mismatches 28; Indels
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                                                                         0;
Qу
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
             11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 30
US-09-248-026-1
; Sequence 1, Application US/09248026
; Patent No. 6093567
; GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
    APPLICANT: A.E.
;
    TITLE OF INVENTION: ADENOVIRUS VECTORS FOR GENE THERAPY
    NUMBER OF SEQUENCES: 10
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: BAKER & BOTTS, L.L.P.
      STREET: 30 ROCKEFELLER PLAZA
      CITY: NEW YORK
      STATE: NEW YORK
      COUNTRY: USA
      ZIP: 10112
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/09/248,026
      FILING DATE: 10-FEB-1999
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 08/895,194
      FILING DATE: 16-JUL-1997
    ATTORNEY/AGENT INFORMATION:
      NAME: Seide, Rochelle K.
      REGISTRATION NUMBER: 32,300
;
      REFERENCE/DOCKET NUMBER: A30668-C
    TELECOMMUNICATION INFORMATION:
     TELEPHONE: (212) 705-5000
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TELEFAX: (212) 705-5020
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
US-09-248-026-1
                        31.6%; Score 32.2; DB 3; Length 6129;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
                              0; Mismatches 28; Indels
 Matches 49; Conservative
                                                             0; Gaps
                                                                         0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                           11 1 1111 11
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 31
US-08-681-838A-1
; Sequence 1, Application US/08681838A
; Patent No. 6245735
; GENERAL INFORMATION:
    APPLICANT: Pier, Gerald B
    TITLE OF INVENTION: Methods and Products for Treating
    TITLE OF INVENTION: Pseudomonas Infection
    NUMBER OF SEQUENCES: 5
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Wolf, Greenfield & Sacks PC
      STREET: 600 Atlantic Avenue
      CITY: Boston
      STATE: MA
      COUNTRY: USA
      ZIP: 02210
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/681,838A
      FILING DATE:
      CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
      NAME: Gates, Edward R
      REGISTRATION NUMBER: 31,616
      REFERENCE/DOCKET NUMBER: B0801/7054
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 617-720-3500
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TELEFAX: 617-720-2441
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
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      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA to mRNA
    HYPOTHETICAL: NO
    ANTI-SENSE: NO
    ORIGINAL SOURCE:
     ORGANISM: Homo sapiens
    FEATURE:
     NAME/KEY: CDS
     LOCATION: 133..4575
US-08-681-838A-1
                       31.6%; Score 32.2; DB 3; Length 6129;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
                           0; Mismatches 28; Indels
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 Matches 49; Conservative
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Qу
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Db
          65 TTGTCACTTTCCGAGGA 81
Qу
            11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 32
US-09-174-077-1
; Sequence 1, Application US/09174077
; Patent No. 6329422
; GENERAL INFORMATION:
; APPLICANT: Fischer, Horst
  APPLICANT: Illek, Beate
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
  FILE REFERENCE: 200116.403C1
; CURRENT APPLICATION NUMBER: US/09/174,077
 CURRENT FILING DATE: 1998-10-16
 EARLIER APPLICATION NUMBER: US 08/951,912
 EARLIER FILING DATE: 1997-10-16
; NUMBER OF SEQ ID NOS: 6
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 6129
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-174-077-1
 Query Match
                       31.6%; Score 32.2; DB 4; Length 6129;
 Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches 49; Conservative 0; Mismatches 28; Indels
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Qу
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Db
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          65 TTGTCACTTTCCGAGGA 81
Qу
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Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 33
US-09-174-077-5
; Sequence 5, Application US/09174077
; Patent No. 6329422
; GENERAL INFORMATION:
; APPLICANT: Fischer, Horst
; APPLICANT: Illek, Beate
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
  FILE REFERENCE: 200116.403C1
  CURRENT APPLICATION NUMBER: US/09/174,077
  CURRENT FILING DATE: 1998-10-16
  EARLIER APPLICATION NUMBER: US 08/951,912
  EARLIER FILING DATE: 1997-10-16
  NUMBER OF SEQ ID NOS: 6
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5
   LENGTH: 6129
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-174-077-5
                        31.6%; Score 32.2; DB 4; Length 6129;
 Query Match
  Best Local Similarity 63.6%; Pred. No. 0.059;
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Qy
             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qy
             1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 34
PCT-US93-11667-1
; Sequence 1, Application PC/TUS9311667
  GENERAL INFORMATION:
    APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith, APPLICANT: A.E.
    TITLE OF INVENTION: GENE THERAPY FOR CYSTIC FIBROSIS
    NUMBER OF SEQUENCES: 9
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: LAHIVE & COCKFIELD
      STREET: 60 STATE STREET, SUITE 510
      CITY: BOSTON
      STATE: MASSACHUSETTS
      COUNTRY: USA
      ZIP: 02109
    COMPUTER READABLE FORM:
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MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: ASCII
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: PCT/US93/11667
      FILING DATE: 02-DEC-1993
      CLASSIFICATION:
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 07/985,478
      FILING DATE: 02-DEC-1992
    ATTORNEY/AGENT INFORMATION:
      NAME: Hanley, Elizabeth A.
      REGISTRATION NUMBER: 33,505
      REFERENCE/DOCKET NUMBER: NZI-014CP2
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 227-7400
      TELEFAX: (617) 227-5941
;
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6129 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
PCT-US93-11667-1
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  Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
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                                                                         0;
          49; Conservative
  Matches
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Qу
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Db
          65 TTGTCACTTTCCGAGGA 81
Qy
             11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 35
US-08-466-886-16
; Sequence 16, Application US/08466886
; Patent No. 5776677
   GENERAL INFORMATION:
    APPLICANT: Tsui, Lap-Chee
    APPLICANT: Riordan, John R.
    APPLICANT: Rommens, Johanna M.
    APPLICANT: Kerem, Bat-Sheva
    APPLICANT: Collins, Francis S.
    APPLICANT: Iannuzzi, Michael C.
    APPLICANT: Drumm, Mitchell L.
    APPLICANT: Buckwald, Manuel
    TITLE OF INVENTION: Cystic Fibrosis Gene
```

```
NUMBER OF SEQUENCES: 43
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX
      STREET: 1100 New York Avenue, N.W.
      CITY: Washington
      STATE: DC
      COUNTRY: USA
      ZIP: 20005
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/466,886
      FILING DATE: 06-JUN-1995
     CLASSIFICATION: 435
    ATTORNEY/AGENT INFORMATION:
      NAME: Goldstein, Jorge A.
      REGISTRATION NUMBER: 29,021
      REFERENCE/DOCKET NUMBER: 1329.0010006
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 202-371-2600
      TELEFAX: 202-371-2540
  INFORMATION FOR SEQ ID NO: 16:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6130 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
US-08-466-886-16
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 Best Local Similarity 63.6%; Pred. No. 0.059;
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                                                             0; Gaps
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           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
QУ
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        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 36
US-08-604-488-1
; Sequence 1, Application US/08604488
; Patent No. 5863770
; GENERAL INFORMATION:
    APPLICANT: TSUI, Lap-Chee
    APPLICANT: ROMMENS, Johanna M.
    TITLE OF INVENTION: Stable Propagation of Modified Full
```

```
TITLE OF INVENTION: Length Cystic Fibrosis Transmembrane Conductance
Regulator
    TITLE OF INVENTION: Protein cDNA in Heterologous Systems
    NUMBER OF SEQUENCES: 1
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Bell, Seltzer, Park & Gibson
      STREET: 1211 East Morehead Street
      CITY: Charlotte
      STATE: No. 5863770th Carolina
      COUNTRY: U.S.A.
      ZIP: 34009
    COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/604,488
      FILING DATE:
      CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: US/08/030,081
      FILING DATE:
    ATTORNEY/AGENT INFORMATION:
     NAME: Layton, Jr., Samuel G
      REGISTRATION NUMBER: 22,807
      REFERENCE/DOCKET NUMBER: 3477-61
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 704-377-1561
      TELEFAX: 704-334-2014
      TELEX: 57-5102
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6130 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    HYPOTHETICAL: NO
    ANTI-SENSE: YES
    ORIGINAL SOURCE:
      TISSUE TYPE: Epithelial
      CELL TYPE: Epithelial cell
    IMMEDIATE SOURCE:
      CLONE: mutant CF gene
    POSITION IN GENOME:
      CHROMOSOME/SEGMENT: 7
      MAP POSITION: XV2C
      UNITS: bp
US-08-604-488-1
                        31.6%; Score 32.2; DB 2; Length 6130;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.059;
  Matches 49; Conservative 0; Mismatches 28; Indels
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Qv
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Db
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          65 TTGTCACTTTCCGAGGA 81
QУ
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        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 37
US-08-469-461-1
; Sequence 1, Application US/08469461B
; Patent No. 5981178
: GENERAL INFORMATION:
; APPLICANT: Tsui, Lap-Chee
; APPLICANT: Rommins, Johanna M.
; APPLICANT: Kerem, Bat-Sheva
; TITLE OF INVENTION: Introns and Exons of the Cystic Fibrosis Gene and
; TITLE OF INVENTION: Mutations at Various Positions of the Gene
; FILE REFERENCE: 3477-61, 033477/139840
  CURRENT APPLICATION NUMBER: US/08/469,461B
  CURRENT FILING DATE: 1995-06-06
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 6130
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)..(4572)
US-08-469-461-1
                        31.6%; Score 32.2; DB 2; Length 6130;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.059;
 Matches 49; Conservative 0; Mismatches 28; Indels
                                                             0; Gaps
                                                                         0;
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Qy
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             1605 TTCTCAGTTTTCCTGGA 1621
RESULT 38
US-07-890-609-1
; Sequence 1, Application US/07890609C
; Patent No. 6001588
; GENERAL INFORMATION:
; APPLICANT: Tsui, Lap-Chee
; APPLICANT: Rommins, Johanna M.
 APPLICANT: Kerem, Bat-Sheva
  TITLE OF INVENTION: Introns and Exons of the Cystic Fibrosis Gene and
 TITLE OF INVENTION: Mutations at Various Positions of the Gene
; FILE REFERENCE: 3477-61, 033477/139840
; CURRENT APPLICATION NUMBER: US/07/890,609C
; CURRENT FILING DATE: 1992-07-13
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NUMBER OF SEQ ID NOS: 33
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 6130
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)..(4572)
US-07-890-609-1
                        31.6%; Score 32.2; DB 3; Length 6130;
 Query Match
                        63.6%; Pred. No. 0.059;
  Best Local Similarity
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 Matches
          49; Conservative
                                               28; Indels
                                                              0; Gaps
                                                                          0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                            Db
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qy
             11 111 111 1 111
Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 39
US-08-030-081-1
; Sequence 1, Application US/08030081
; Patent No. 6063913
  GENERAL INFORMATION:
    APPLICANT: TSUI, Lap-Chee
    APPLICANT: ROMMENS, Johanna M.
    TITLE OF INVENTION: Stable Propagation of Modified Full
    TITLE OF INVENTION: Length Cystic Fibrosis Transmembrane Conductance
Regulator
    TITLE OF INVENTION: Protein cDNA in Heterologous Systems
    NUMBER OF SEQUENCES: 1
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Bell, Seltzer, Park & Gibson
      STREET: 1211 East Morehead Street
      CITY: Charlotte
      STATE: No. 6063913th Carolina
      COUNTRY: U.S.A.
      ZIP: 34009
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/030,081
      FILING DATE: 19930412
      CLASSIFICATION: 435
    ATTORNEY/AGENT INFORMATION:
      NAME: Layton, Jr., Samuel G
      REGISTRATION NUMBER: 22,807
      REFERENCE/DOCKET NUMBER: 3477-61
    TELECOMMUNICATION INFORMATION:
```

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TELEPHONE: 704-377-1561
      TELEFAX: 704-334-2014
      TELEX: 57-5102
  INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6130 base pairs
      TYPE: NUCLEIC ACID
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    HYPOTHETICAL: NO
    ANTI-SENSE: YES
    ORIGINAL SOURCE:
      TISSUE TYPE: Epithelial
      CELL TYPE: Epithelial cell
    IMMEDIATE SOURCE:
      CLONE: mutant CF gene
    POSITION IN GENOME:
      CHROMOSOME/SEGMENT:
      MAP POSITION: XV2C
      UNITS: bp
US-08-030-081-1
 Query Match
                        31.6%; Score 32.2; DB 3; Length 6130;
 Best Local Similarity 63.6%; Pred. No. 0.059;
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 Matches 49; Conservative
                                               28; Indels
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                                                                 Gaps
                                                                        0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
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Db
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RESULT 40
US-08-469-617-16
; Sequence 16, Application US/08469617
; Patent No. 6201107
  GENERAL INFORMATION:
    APPLICANT: Tsui, Lap-Chee
    APPLICANT: Riordan, John R.
    APPLICANT: Rommens, Johanna M.
    APPLICANT: Kerem, Bat-Sheva
    APPLICANT: Collins, Francis S.
    APPLICANT: Iannuzzi, Michael C.
    APPLICANT: Drumm, Mitchell L.
    APPLICANT: Buckwald, Manuel
    TITLE OF INVENTION: Cystic Fibrosis Gene
    NUMBER OF SEQUENCES: 43
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
      STREET: 1100 New York Avenue, N.W.
      CITY: Washington
      STATE: DC
      COUNTRY: USA
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ZIP: 20005
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/469,617
      FILING DATE: 06-JUN-1995
      CLASSIFICATION: 800
    ATTORNEY/AGENT INFORMATION:
      NAME: Goldstein, Jorge A.
      REGISTRATION NUMBER: 29,021
      REFERENCE/DOCKET NUMBER: 1329.0010008
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 202-371-2600
      TELEFAX: 202-371-2540
   INFORMATION FOR SEQ ID NO: 16:
    SEQUENCE CHARACTERISTICS:
    LENGTH: 6130 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    FEATURE:
      NAME/KEY: CDS
      LOCATION: 133..4572
US-08-469-617-16
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        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 41
5240846-4
; Patent No. 5240846
    APPLICANT: Collins, Francis S.; Wilson, James C.
    TITLE OF INVENTION: GENE THERAPY VECTOR FOR CYSTIC
; FIBROSIS
    NUMBER OF SEQUENCES: 5
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/584,275
      FILING DATE: 18-SEP-1990
    PRIOR APPLICATION DATA:
    APPLICATION NUMBER: 399,945
     FILING DATE: 24-AUG-1989
     APPLICATION NUMBER: 401,609
     FILING DATE: 31-AUG-1989
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Qy
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            Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 42
US-08-793-618-1
; Sequence 1, Application US/08793618
; Patent No. 6265218
; GENERAL INFORMATION:
    APPLICANT: SEEBER, Stefan
    TITLE OF INVENTION: GENE THERAPY METHOD USING DNA VECTORS
    TITLE OF INVENTION: WITHOUT A SELECTION MARKER GENE
    NUMBER OF SEQUENCES: 5
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Felfe & Lynch
      STREET: 805 Third Avenue
      CITY: New York City
      STATE: New York
      COUNTRY: USA
     ZIP: 10022
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Diskette, 3.50 inch, 1.44mb
      COMPUTER: IBM PS/2
      OPERATING SYSTEM: PC-DOS
      SOFTWARE: Wordperfect
    CURRENT APPLICATION DATA:
    APPLICATION NUMBER: US/08/793,618
     FILING DATE: June 10, 1997
      CLASSIFICATION: 514
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: PCT/EP95/03027
      FILING DATE: July 31, 1995
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: DE P 44 28 402.0
      FILING DATE: 11-AUG-1994
    ATTORNEY/AGENT INFORMATION:
      NAME: Susan L. Hess
      REGISTRATION NUMBER: 37,350
      REFERENCE/DOCKET NUMBER: BOER 1075 PCT
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (212) 688-9200
      TELEFAX: (212) 838-3884
 INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
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LENGTH: 8225 base pairs
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RESULT 43
US-09-794-431-1
; Sequence 1, Application US/09794431
; Patent No. 6573100
   GENERAL INFORMATION:
        APPLICANT: SEEBER, Stefan
        TITLE OF INVENTION: GENE THERAPY METHOD USING DNA VECTORS
                           WITHOUT A SELECTION MARKER GENE
        NUMBER OF SEQUENCES: 5
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: Felfe & Lynch
             STREET: 805 Third Avenue
             CITY: New York City
             STATE: New York
             COUNTRY: USA
             ZIP: 10022
        COMPUTER READABLE FORM:
             MEDIUM TYPE: Diskette, 3.50 inch, 1.44mb
             COMPUTER: IBM PS/2
             OPERATING SYSTEM: PC-DOS
             SOFTWARE: Wordperfect
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/09/794,431
             FILING DATE: 27-Feb-2001
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: 08/793,618
             FILING DATE: <Unknown>
             APPLICATION NUMBER: DE P 44 28 402.0
             FILING DATE: 11-AUG-1994
        ATTORNEY/AGENT INFORMATION:
             NAME: Susan L. Hess
             REGISTRATION NUMBER: 37,350
             REFERENCE/DOCKET NUMBER: BOER 1075 PCT
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: (212) 688-9200
             TELEFAX: (212) 838-3884
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INFORMATION FOR SEQ ID NO: 1:
         SEQUENCE CHARACTERISTICS:
             LENGTH: 8225 base pairs
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             TOPOLOGY: linear
        MOLECULE TYPE: cDNA
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Qу
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        2272 TTCTCAGTTTTCCTGGA 2288
RESULT 44
US-08-836-022A-3/c
; Sequence 3, Application US/08836022A
; Patent No. 6001557
  GENERAL INFORMATION:
    APPLICANT: Trustees of the University of Pennsylvania
    APPLICANT: Wilson, James M. APPLICANT: Fisher, Krishna J.
    APPLICANT: Chen, Shu-Jen
;
    APPLICANT: Weitzman, Matthew
    TITLE OF INVENTION: Improved Adenovirus Virus and
    NUMBER OF SEQUENCES: 10
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Howson and Howson
      STREET: Spring House Corporate Cntr, P O Box 457
      CITY: Spring House
      STATE: Pennsylvania
;
      COUNTRY: USA
;
      ZIP: 19477
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
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    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/836,022A
      FILING DATE:
      CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 08/331,381
      FILING DATE: 28-OCT-1994
    ATTORNEY/AGENT INFORMATION:
      NAME: Bak, Mary E.
      REGISTRATION NUMBER: 31,215
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REFERENCE/DOCKET NUMBER: GNVPN.008PCT
     TELECOMMUNICATION INFORMATION:
      TELEPHONE: 215-540-9200
       TELEFAX: 215-540-5818
   INFORMATION FOR SEQ ID NO: 3:
     SEQUENCE CHARACTERISTICS:
      LENGTH: 9972 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: double
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  Best Local Similarity
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        7089 TTCTCAGTTTTCCTGGA 7073
RESULT 45
US-09-427-048A-3/c
; Sequence 3, Application US/09427048A
 Patent No. 6203975
   GENERAL INFORMATION:
        APPLICANT: Trustees of the University of Pennsylvania
                   Wilson, James M.
                   Fisher, Krishna J.
                   Chen, Shu-Jen
                   Weitzman, Matthew
        TITLE OF INVENTION: Improved Adenovirus Virus and
                           Methods of Use Thereof
        NUMBER OF SEQUENCES: 10
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: Howson and Howson
             STREET: Spring House Corporate Cntr, P O Box 457
             CITY: Spring House
             STATE: Pennsylvania
             COUNTRY: USA
             ZIP: 19477
        COMPUTER READABLE FORM:
             MEDIUM TYPE: Floppy disk
             COMPUTER: IBM PC compatible
             OPERATING SYSTEM: PC-DOS/MS-DOS
             SOFTWARE: PatentIn Release #1.0, Version #1.30
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/09/427,048A
             FILING DATE: 21-Oct-1999
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: 08/836,022
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FILING DATE: <Unknown>
        ATTORNEY/AGENT INFORMATION:
             NAME: Bak, Mary E.
             REGISTRATION NUMBER: 31,215
             REFERENCE/DOCKET NUMBER: GNVPN.008PCT
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: 215-540-9200
             TELEFAX: 215-540-5818
    INFORMATION FOR SEO ID NO: 3:
        SEQUENCE CHARACTERISTICS:
             LENGTH: 9972 base pairs
             TYPE: nucleic acid
             STRANDEDNESS: double
             TOPOLOGY: unknown
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US-09-423-744A-1
; Sequence 1, Application US/09423744A
; Patent No. 6372500
   GENERAL INFORMATION:
        APPLICANT: HSC Research and Development Limited Partnership
        TITLE OF INVENTION: Episomal Expression Cassettes for Gene
                            Therapy
        NUMBER OF SEQUENCES: 19
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: Rockey, Milnamov & Katz, Ltd.
             STREET: 180 N. Stetson Avenue, Suite 4700
             CITY: Chicago
             STATE: Illinois
             COUNTRY: USA
             ZIP: 60601
        COMPUTER READABLE FORM:
             MEDIUM TYPE: Floppy disk
             COMPUTER: IBM PC compatible
             OPERATING SYSTEM: PC-DOS/MS-DOS
             SOFTWARE: PatentIn Release #1.0, Version #1.30
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/09/423,744A
             FILING DATE: 12-No. 6372500-1999
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
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APPLICATION NUMBER: PCT/CA98/00478
          FILING DATE: May 14, 1998
    ATTORNEY/AGENT INFORMATION:
          NAME: Lisa V. Mueller
          REFERENCE/DOCKET NUMBER: DWW6064P0020US
INFORMATION FOR SEQ ID NO: 1:
    SEQUENCE CHARACTERISTICS:
          LENGTH: 12143 base pairs
          TYPE: nucleic acid
          STRANDEDNESS: double
          TOPOLOGY: circular
    MOLECULE TYPE: other nucleic acid
          DESCRIPTION: /desc = "Mixture of genomic DNA,
    FEATURE:
         NAME/KEY: enhancer
          LOCATION: 8..2570
          IDENTIFICATION METHOD:
          OTHER INFORMATION: /standard_name= "K18
          Enhancer/Promoter"
          /note= "DNA fragment was obtained by PCR-cloning and minor
         modifications were introduced for the purpose of PCR."
    FEATURE:
         NAME/KEY: intron
          LOCATION: 2571..3318
          IDENTIFICATION METHOD:
          OTHER INFORMATION: /standard name= "K18 intron 1"
          /note= "DNA fragment was obtained by PCR-cloning and
         modifications were introduced to improve the splicing
          efficiency."
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          virus translational enhancer"
          /note= "Fragment was synthesized chemically."
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          LOCATION: 3355..7948
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          OTHER INFORMATION: /standard name= "CFTR cDNA"
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          NAME/KEY: misc feature
          LOCATION: 7949..7984
          IDENTIFICATION METHOD:
          OTHER INFORMATION: /standard_name= "pBluescript II
          KS(+) multiple cloning site"
    FEATURE:
          NAME/KEY: intron
          LOCATION: 8507..8572
          IDENTIFICATION METHOD:
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          LOCATION: 9178..9212
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OTHER INFORMATION: /standard name= "SV40"
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RESULT 47
US-08-469-461-3
; Sequence 3, Application US/08469461B
; Patent No. 5981178
; GENERAL INFORMATION:
; APPLICANT: Tsui, Lap-Chee
; APPLICANT: Rommins, Johanna M.
 APPLICANT: Kerem, Bat-Sheva
 TITLE OF INVENTION: Introns and Exons of the Cystic Fibrosis Gene and
  TITLE OF INVENTION: Mutations at Various Positions of the Gene
  FILE REFERENCE: 3477-61, 033477/139840
; CURRENT APPLICATION NUMBER: US/08/469,461B
; CURRENT FILING DATE: 1995-06-06
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.0
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   ORGANISM: Homo sapiens
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RESULT 48
US-07-890-609-3
; Sequence 3, Application US/07890609C
; Patent No. 6001588
; GENERAL INFORMATION:
 APPLICANT: Tsui, Lap-Chee
 APPLICANT: Rommins, Johanna M.
; APPLICANT: Kerem, Bat-Sheva
; TITLE OF INVENTION: Introns and Exons of the Cystic Fibrosis Gene and
; TITLE OF INVENTION: Mutations at Various Positions of the Gene
; FILE REFERENCE: 3477-61, 033477/139840
; CURRENT APPLICATION NUMBER: US/07/890,609C
; CURRENT FILING DATE: 1992-07-13
; NUMBER OF SEQ ID NOS: 33
 SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
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   ORGANISM: Homo sapiens
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 Best Local Similarity 63.6%; Pred. No. 0.099;
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RESULT 49
US-09-252-991A-1019/c
; Sequence 1019, Application US/09252991A
; Patent No. 6551795
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; GENERAL INFORMATION:
 APPLICANT: Marc J. Rubenfield et al.
  TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
  CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
  PRIOR APPLICATION NUMBER: US 60/094,190
  PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 1019
   LENGTH: 1251
   TYPE: DNA
   ORGANISM: Pseudomonas aeruginosa
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          63 TGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGGAGGC 101
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             1007 TGTCGACGATGGTCGAGTCGATCACCCAGCCCTTGGGGC 969
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US-09-252-991A-1036
; Sequence 1036, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
 APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
 CURRENT FILING DATE: 1999-02-18
  PRIOR APPLICATION NUMBER: US 60/074,788
  PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
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   LENGTH: 2847
   TYPE: DNA
   ORGANISM: Pseudomonas aeruginosa
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 Best Local Similarity 56.6%; Pred. No. 0.23;
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OM nucleic - nucleic search, using sw model

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April 29, 2004, 17:06:46; Search time 99.1938 Seconds

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Perfect score: 102

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Searched:

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Listing first 50 summaries

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- 17: /cgn2 6/ptodata/2/pubpna/US10 NEW PUB.seq:*
- /cgn2 6/ptodata/2/pubpna/US60 NEW PUB.seq:*
- /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:* 19:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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Result

Query

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	2	102	100.0	6043	10	US-09-989-981A-9	Sequence 9, Appli
С	3	87.6	85.9	2669	10	US-09-989-981A-7	Sequence 7, Appli
	4	32.2	31.6	180	9	US-09-864-761-27920	Sequence 27920, A
	5	32.2	31.6	240	12	US-10-441-643-1	Sequence 1, Appli
	6	32.2	31.6	420	9	US-09-756-095-64	Sequence 64, Appl
	7	32.2	31.6	420	10	US-09-941-492-64	Sequence 64, Appl
	8	32.2	31.6	420	10	US-09-756-096A-64	Sequence 64, Appl
	9	32.2	31.6	420	10	US-09-838-858-64	Sequence 64, Appl
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; Publication No. US20030049730A1
; GENERAL INFORMATION:
  APPLICANT: Hobbs, Helen H.
  APPLICANT: Shan, Bei
  APPLICANT: Barnes, Robert
  APPLICANT: Tian, Hui
  APPLICANT: Tularik Inc.
  APPLICANT: Board of Regents, The University of Texas System
  TITLE OF INVENTION: ABCG5 and ABCG8: Compositions and Methods of Use
  FILE REFERENCE: 018781-007320US
  CURRENT APPLICATION NUMBER: US/09/989,981A
  CURRENT FILING DATE: 2002-07-23
  PRIOR APPLICATION NUMBER: US 60/252,235
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/253,645
  PRIOR FILING DATE: 2000-11-28
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; Publication No. US20030049730A1
; GENERAL INFORMATION:
; APPLICANT: Hobbs, Helen H.
 APPLICANT: Shan, Bei
  APPLICANT: Barnes, Robert
; APPLICANT: Tian, Hui
; APPLICANT: Tularik Inc.
; APPLICANT: Board of Regents, The University of Texas System
; TITLE OF INVENTION: ABCG5 and ABCG8: Compositions and Methods of Use
; FILE REFERENCE: 018781-007320US
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CURRENT APPLICATION NUMBER: US/09/989,981A
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  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/253,645
  PRIOR FILING DATE: 2000-11-28
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; Publication No. US20030049730A1
; GENERAL INFORMATION:
  APPLICANT: Hobbs, Helen H.
; APPLICANT: Shan, Bei
; APPLICANT: Barnes, Robert
; APPLICANT: Tian, Hui
  APPLICANT: Tularik Inc.
  APPLICANT: Board of Regents, The University of Texas System
  TITLE OF INVENTION: ABCG5 and ABCG8: Compositions and Methods of Use
  FILE REFERENCE: 018781-007320US
  CURRENT APPLICATION NUMBER: US/09/989,981A
  CURRENT FILING DATE: 2002-07-23
  PRIOR APPLICATION NUMBER: US 60/252,235
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/253,645
  PRIOR FILING DATE: 2000-11-28
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; Sequence 27920, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
; APPLICANT: Rank, David R.
 APPLICANT: Hanzel, David K.
  APPLICANT: Chen, Wensheng
  TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES
USEFUL FOR
; TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
  FILE REFERENCE: Aeomica-X-1
  CURRENT APPLICATION NUMBER: US/09/864,761
  CURRENT FILING DATE: 2001-05-23
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  PRIOR FILING DATE: 2000-02-04
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  PRIOR APPLICATION NUMBER: PCT/US01/00668
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  PRIOR APPLICATION NUMBER: PCT/US01/00662
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; Publication No. US20040072208A1
; GENERAL INFORMATION:
  APPLICANT: Warthoe, Peter
  TITLE OF INVENTION: Surface Acoustic Wave Sensors and Method for Detecting
Target
  TITLE OF INVENTION: Analytes
  FILE REFERENCE: A-71523
; CURRENT APPLICATION NUMBER: US/10/441,643
; CURRENT FILING DATE: 2003-05-20
; PRIOR APPLICATION NUMBER: US 60/383,247
; PRIOR FILING DATE: 2002-05-23
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; Patent No. US20020115207A1
; GENERAL INFORMATION:
  APPLICANT: Mitchell, Lloyd G.
  APPLICANT: Garcia-Blanco, Mariano A.
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-B-A 072874.0134
  CURRENT APPLICATION NUMBER: US/09/756,095
   CURRENT FILING DATE: 2001-01-08
   PRIOR APPLICATION NUMBER: 09/158,863
   PRIOR FILING DATE: 1998-09-23
   PRIOR APPLICATION NUMBER: 09/133,717
   PRIOR FILING DATE: 1998-08-13
   PRIOR APPLICATION NUMBER: 09/087,233
   PRIOR FILING DATE: 1998-05-28
  PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
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   PRIOR FILING DATE: 1995-12-07
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    OTHER INFORMATION: transmembrane regulator-derived sequences and His
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; Publication No. US20030027250A1
; GENERAL INFORMATION:
  APPLICANT: Mitchell, Lloyd
  APPLICANT: Garcia-Blanco, Mariano M.
  APPLICANT: Puttaraju, Madaiah
  APPLICANT: Mansfield, Gary S.
  TITLE OF INVENTION: METHODS OF COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-BAE (072874.0156)
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  CURRENT FILING DATE: 2002-04-01
   PRIOR APPLICATION NUMBER: 09/838,858
  PRIOR FILING DATE: 2001-04-20
  PRIOR APPLICATION NUMBER: 09/756,096
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  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
   PRIOR APPLICATION NUMBER: 09/087,233
   PRIOR FILING DATE: 1998-05-28
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   OTHER INFORMATION: Trans-spliced product comprising cystic fibrosis
   OTHER INFORMATION: transmembrane regulator-derived sequences and His
    OTHER INFORMATION: tag sequences
US-09-941-492-64
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QУ
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; Publication No. US20030077754A1
; GENERAL INFORMATION:
; APPLICANT: Mitchell, Lloyd G.
; APPLICANT: Garcia-Blanco, Mariano A.
; APPLICANT: Puttaraju, Madaiah
  APPLICANT: Mansfield, Gary S.
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-B-A-B 072874.0135
  CURRENT APPLICATION NUMBER: US/09/756,096A
  CURRENT FILING DATE: 2001-01-08
  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
   PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
   PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
  PRIOR APPLICATION NUMBER: 60/008,317
  PRIOR FILING DATE: 1995-12-15
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    OTHER INFORMATION: transmembrane regulator-derived sequences and His
    OTHER INFORMATION: tag sequence
US-09-756-096A-64
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Qу
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US-09-838-858-64

; Sequence 64, Application US/09838858

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; Publication No. US20030148937A1
; GENERAL INFORMATION:
  APPLICANT: Mansfield, Gary S.
  APPLICANT: Mitchell, Lloyd G.
  APPLICANT: Garcia-Blanco, Mariano A.
  APPLICANT: Walsh, Christopher E.
  APPLICANT: Chao, Hengjun
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-BAD 072874.01
  CURRENT APPLICATION NUMBER: US/09/838,858
  CURRENT FILING DATE: 2001-04-20
  PRIOR APPLICATION NUMBER: 09/756,096
  PRIOR FILING DATE: 2001-02-08
  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
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   OTHER INFORMATION: His-tag sequence
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; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
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APPLICANT: Chen, Wensheng
  TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES
USEFUL FOR
  TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
  FILE REFERENCE: Aeomica-X-1
  CURRENT APPLICATION NUMBER: US/09/864,761
  CURRENT FILING DATE: 2001-05-23
  PRIOR APPLICATION NUMBER: US 60/180,312
  PRIOR FILING DATE: 2000-02-04
  PRIOR APPLICATION NUMBER: US 60/207,456
  PRIOR FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/632,366
  PRIOR FILING DATE: 2000-08-03
  PRIOR APPLICATION NUMBER: GB 24263.6
  PRIOR FILING DATE: 2000-10-04
  PRIOR APPLICATION NUMBER: US 60/236,359
  PRIOR FILING DATE: 2000-09-27
  PRIOR APPLICATION NUMBER: PCT/US01/00666
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00667
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00664
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00669
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00665
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: PCT/US01/00663
  PRIOR FILING DATE: 2001-01-30
   PRIOR APPLICATION NUMBER: PCT/US01/00662
   PRIOR FILING DATE: 2001-01-30
   PRIOR APPLICATION NUMBER: PCT/US01/00661
   PRIOR FILING DATE: 2001-01-30
   PRIOR APPLICATION NUMBER: PCT/US01/00670
  PRIOR FILING DATE: 2001-01-30
  PRIOR APPLICATION NUMBER: US 60/234,687
  PRIOR FILING DATE: 2000-09-21
  PRIOR APPLICATION NUMBER: US 09/608,408
   PRIOR FILING DATE: 2000-06-30
   PRIOR APPLICATION NUMBER: US 09/774,203
   PRIOR FILING DATE: 2001-01-29
   NUMBER OF SEQ ID NOS: 49117
   SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 11433
    LENGTH: 494
    TYPE: DNA
    ORGANISM: Homo sapiens
    FEATURE:
    OTHER INFORMATION: MAP TO AC000111.1
    OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.92
    OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 0.94
    OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 0.94
    OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.1
    OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL =
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OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.1

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31.6%; Score 32.2; DB 9; Length 494;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.14;
                              0; Mismatches
                                                           0; Gaps
                                                                      0;
 Matches
          49; Conservative
                                             28; Indels
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
QУ
            280 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 339
Db
         65 TTGTCACTTTCCGAGGA 81
Qу
             Db
         340 TTCTCAGTTTTCCTGGA 356
RESULT 11
US-10-300-683-247
; Sequence 247, Application US/10300683
; Publication No. US20030235834A1
; GENERAL INFORMATION:
  APPLICANT: Dunlop, Charles L.M.
  APPLICANT: Weisel, James M.
  TITLE OF INVENTION: APPROACHES TO IDENTIFY CYSTIC FIBROSIS
  FILE REFERENCE: CHARDUN.010A
  CURRENT APPLICATION NUMBER: US/10/300,683
  CURRENT FILING DATE: 2002-11-19
  PRIOR APPLICATION NUMBER: 60/333,531
  PRIOR FILING DATE: 2001-11-19
  NUMBER OF SEQ ID NOS: 554
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEO ID NO 247
   LENGTH: 831
   TYPE: DNA
   ORGANISM: Artificial Sequence
   OTHER INFORMATION: Diagnostic Oligonucleotide
US-10-300-683-247
                       31.6%;
                               Score 32.2; DB 16; Length 831;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.16;
          49; Conservative
                              0; Mismatches
                                             28;
                                                  Indels
                                                               Gaps
                                                                      0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             328 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 387
Db
          65 TTGTCACTTTCCGAGGA 81
Qу
             388 TTCTCAGTTTTCCTGGA 404
Db
RESULT 12
US-09-756-095-105
; Sequence 105, Application US/09756095
; Patent No. US20020115207A1
; GENERAL INFORMATION:
; APPLICANT: Mitchell, Lloyd G.
```

```
APPLICANT: Garcia-Blanco, Mariano A.
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-B-A 072874.0134
  CURRENT APPLICATION NUMBER: US/09/756,095
  CURRENT FILING DATE: 2001-01-08
  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
 PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
  PRIOR APPLICATION NUMBER: 60/008,317
  PRIOR FILING DATE: 1995-12-07
 NUMBER OF SEQ ID NOS: 105
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 105
   LENGTH: 3069
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: CFTR PTM sequence
US-09-756-095-105
                         31.6%; Score 32.2; DB 9; Length 3069;
 Query Match
                        63.6%; Pred. No. 0.23;
 Best Local Similarity
 Matches 49; Conservative
                               0; Mismatches
                                                28; Indels
                                                              0; Gaps
                                                                          0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                            21 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 80
Db
          65 TTGTCACTTTCCGAGGA 81
Qy
             Db
          81 TTCTCAGTTTTCCTGGA 97
RESULT 13
US-09-941-492-105
; Sequence 105, Application US/09941492
; Publication No. US20030027250A1
; GENERAL INFORMATION:
; APPLICANT: Mitchell, Lloyd
; APPLICANT: Garcia-Blanco, Mariano M.
; APPLICANT: Puttaraju, Madaiah
; APPLICANT: Mansfield, Gary S.
  TITLE OF INVENTION: METHODS OF COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-BAE (072874.0156)
  CURRENT APPLICATION NUMBER: US/09/941,492
  CURRENT FILING DATE: 2002-04-01
  PRIOR APPLICATION NUMBER: 09/838,858
; PRIOR FILING DATE: 2001-04-20
; PRIOR APPLICATION NUMBER: 09/756,096
  PRIOR FILING DATE: 2001-01-08
```

```
PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
  PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
  NUMBER OF SEQ ID NOS: 125
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 105
   LENGTH: 3069
   TYPE: DNA
   ORGANISM: Artificial Sequence
   OTHER INFORMATION: CFTR PTM sequence
US-09-941-492-105
 Query Match
                         31.6%; Score 32.2; DB 10; Length 3069;
 Best Local Similarity 63.6%; Pred. No. 0.23;
          49; Conservative
                             0; Mismatches
                                                28;
 Matches
                                                     Indels
                                                              0; Gaps
                                                                          0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             21 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 80
Db
Qу
          65 TTGTCACTTTCCGAGGA 81
             81 TTCTCAGTTTTCCTGGA 97
Db
RESULT 14
US-09-756-096A-105
; Sequence 105, Application US/09756096A
; Publication No. US20030077754A1
; GENERAL INFORMATION:
; APPLICANT: Mitchell, Lloyd G.
 APPLICANT: Garcia-Blanco, Mariano A.
  APPLICANT: Puttaraju, Madaiah
  APPLICANT: Mansfield, Gary S.
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-B-A-B 072874.0135
  CURRENT APPLICATION NUMBER: US/09/756,096A
  CURRENT FILING DATE: 2001-01-08
  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
  PRIOR APPLICATION NUMBER: 08/766,354
  PRIOR FILING DATE: 1996-12-13
  PRIOR APPLICATION NUMBER: 60/008,317
  PRIOR FILING DATE: 1995-12-15
  NUMBER OF SEQ ID NOS: 105
  SOFTWARE: FastSEQ for Windows Version 4.0
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; SEQ ID NO 105
   LENGTH: 3069
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: CFTR PTM sequence
US-09-756-096A-105
                         31.6%; Score 32.2; DB 10; Length 3069;
 Query Match
                        63.6%; Pred. No. 0.23;
  Best Local Similarity
          49: Conservative
                               0; Mismatches
                                                28;
                                                    Indels
                                                                  Gaps
                                                                          0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
                           21 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 80
          65 TTGTCACTTTCCGAGGA 81
Qу
             81 TTCTCAGTTTTCCTGGA 97
RESULT 15
US-09-838-858-105
; Sequence 105, Application US/09838858
; Publication No. US20030148937A1
; GENERAL INFORMATION:
  APPLICANT: Mansfield, Gary S.
  APPLICANT: Mitchell, Lloyd G.
 APPLICANT: Garcia-Blanco, Mariano A.
; APPLICANT: Walsh, Christopher E.
 APPLICANT: Chao, Hengjun
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR USE IN
  TITLE OF INVENTION: SPLICEOSOME MEDIATED RNA TRANS-SPLICING
  FILE REFERENCE: A31304-BAD 072874.01
  CURRENT APPLICATION NUMBER: US/09/838,858
  CURRENT FILING DATE: 2001-04-20
  PRIOR APPLICATION NUMBER: 09/756,096
  PRIOR FILING DATE: 2001-02-08
  PRIOR APPLICATION NUMBER: 09/158,863
  PRIOR FILING DATE: 1998-09-23
  PRIOR APPLICATION NUMBER: 09/133,717
  PRIOR FILING DATE: 1998-08-13
  PRIOR APPLICATION NUMBER: 09/087,233
  PRIOR FILING DATE: 1998-05-28
; PRIOR APPLICATION NUMBER: 08/766,354
; PRIOR FILING DATE: 1996-12-13
; PRIOR APPLICATION NUMBER: 60/008,317
; PRIOR FILING DATE: 1995-12-15
  NUMBER OF SEQ ID NOS: 113
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 105
   LENGTH: 3069
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: CFTR PTM sequence
US-09-838-858-105
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31.6%; Score 32.2; DB 10; Length 3069;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.23;
 Matches 49; Conservative 0; Mismatches 28;
                                                  Indels
                                                                      0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            Db
          21 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 80
          65 TTGTCACTTTCCGAGGA 81
Qy
            11 111 111 1 111
          81 TTCTCAGTTTTCCTGGA 97
Db
RESULT 16
US-10-367-507-1
; Sequence 1, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
; APPLICANT: Welsh, Michael J.
 APPLICANT: Ostedgaard, Lynda S.
; APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
  CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
  PRIOR FILING DATE: 2002-02-19
; NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
   LENGTH: 4191
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4191)
US-10-367-507-1
 Query Match
                       31.6%; Score 32.2; DB 16; Length 4191;
 Best Local Similarity 63.6%; Pred. No. 0.25;
 Matches 49; Conservative
                             0; Mismatches
                                             28;
                                                           0; Gaps
                                                                      0;
                                                  Indels
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            D\mathbf{b}
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
QУ
            1605 TTCTCAGTTTTCCTGGA 1621
RESULT 17
US-10-367-507-8
; Sequence 8, Application US/10367507
```

; Publication No. US20030235885A1

```
; GENERAL INFORMATION:
; APPLICANT: Welsh, Michael J.
  APPLICANT: Ostedgaard, Lynda S.
  APPLICANT: Zabner, Joseph
; TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
; TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
  CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
  PRIOR FILING DATE: 2002-02-19
  NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 8
   LENGTH: 4311
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4311)
US-10-367-507-8
 Query Match
                        31.6%; Score 32.2; DB 16; Length 4311;
 Best Local Similarity 63.6%; Pred. No. 0.25;
         49; Conservative 0; Mismatches
 Matches
                                               28; Indels
                                                              0; Gaps
                                                                         0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
             11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 18
US-10-367-507-6
; Sequence 6, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
 APPLICANT: Welsh, Michael J.
 APPLICANT: Ostedgaard, Lynda S.
  APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
  CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
  PRIOR FILING DATE: 2002-02-19
  NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 6
   LENGTH: 4347
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
```

```
NAME/KEY: CDS
  LOCATION: (133)...(4347)
US-10-367-507-6
 Query Match
                      31.6%; Score 32.2; DB 16; Length 4347;
 Best Local Similarity 63.6%; Pred. No. 0.25;
 Matches 49; Conservative
                             0; Mismatches
                                             28: Indels
                                                                     0;
                                                          0:
                                                             Gaps
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
         65 TTGTCACTTTCCGAGGA 81
Qу
            1605 TTCTCAGTTTTCCTGGA 1621
RESULT 19
US-10-367-507-7
; Sequence 7, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
; APPLICANT: Welsh, Michael J.
 APPLICANT: Ostedgaard, Lynda S.
  APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
; CURRENT APPLICATION NUMBER: US/10/367,507
; CURRENT FILING DATE: 2003-02-14
; PRIOR APPLICATION NUMBER: 60/358,074
; PRIOR FILING DATE: 2002-02-19
; NUMBER OF SEQ ID NOS: 16
 SOFTWARE: FastSEQ for Windows Version 4.0
; SEO ID NO 7
   LENGTH: 4347
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4347)
US-10-367-507-7
 Query Match
                       31.6%; Score 32.2; DB 16; Length 4347;
 Best Local Similarity 63.6%; Pred. No. 0.25;
 Matches 49; Conservative 0; Mismatches 28; Indels
                                                          0; Gaps
                                                                     0;
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                         11 | | | | | | | |
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Dh
          65 TTGTCACTTTCCGAGGA 81
Qy
            1605 TTCTCAGTTTTCCTGGA 1621
Db
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US-10-367-507-5
; Sequence 5, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
; APPLICANT: Welsh, Michael J.
 APPLICANT: Ostedgaard, Lynda S.
  APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
; CURRENT FILING DATE: 2003-02-14
; PRIOR APPLICATION NUMBER: 60/358,074
; PRIOR FILING DATE: 2002-02-19
; NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 5
   LENGTH: 4368
   TYPE: DNA
;
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4368)
US-10-367-507-5
                        31.6%; Score 32.2; DB 16; Length 4368;
  Query Match
  Best Local Similarity 63.6%; Pred. No. 0.25;
 Matches 49; Conservative 0; Mismatches
                                               28; Indels
                                                             0; Gaps
                                                                        0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qv
             1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qy
             Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 21
US-10-367-507-4
; Sequence 4, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
  APPLICANT: Welsh, Michael J.
  APPLICANT: Ostedgaard, Lynda S.
; APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
  CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
  PRIOR FILING DATE: 2002-02-19
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEO ID NO 4
  LENGTH: 4371
```

```
TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4371)
US-10-367-507-4
  Query Match
                        31.6%; Score 32.2; DB 16; Length 4371;
  Best Local Similarity 63.6%; Pred. No. 0.25;
                             0; Mismatches
 Matches 49; Conservative
                                            28; Indels
                                                            0; Gaps
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             Db
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Qу
          65 TTGTCACTTTCCGAGGA 81
             11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 22
US-10-367-507-3
; Sequence 3, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
; APPLICANT: Welsh, Michael J.
; APPLICANT: Ostedgaard, Lynda S. ; APPLICANT: Zabner, Joseph
; TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
; TITLE OF INVENTION: AND USES THEREOF
; FILE REFERENCE: AP35027 (072419.0117)
; CURRENT APPLICATION NUMBER: US/10/367.507
; CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
; PRIOR FILING DATE: 2002-02-19
  NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 3
  LENGTH: 4410
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4410)
US-10-367-507-3
  Query Match
                        31.6%; Score 32.2; DB 16; Length 4410;
  Best Local Similarity 63.6%; Pred. No. 0.25;
 Matches
          49; Conservative 0; Mismatches 28; Indels
                                                           0; Gaps
                                                                      0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
             1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
Qу
             Db
        1605 TTCTCAGTTTTCCTGGA 1621
```

```
RESULT 23
US-10-367-507-2
; Sequence 2, Application US/10367507
; Publication No. US20030235885A1
; GENERAL INFORMATION:
  APPLICANT: Welsh, Michael J.
  APPLICANT: Ostedgaard, Lynda S.
  APPLICANT: Zabner, Joseph
  TITLE OF INVENTION: CFTR WITH A PARTIALLY DELETED R DOMAIN
  TITLE OF INVENTION: AND USES THEREOF
  FILE REFERENCE: AP35027 (072419.0117)
  CURRENT APPLICATION NUMBER: US/10/367,507
  CURRENT FILING DATE: 2003-02-14
  PRIOR APPLICATION NUMBER: 60/358,074
  PRIOR FILING DATE: 2002-02-19
  NUMBER OF SEQ ID NOS: 16
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 2
   LENGTH: 4419
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (133)...(4419)
US-10-367-507-2
  Query Match
                         31.6%; Score 32.2; DB 16; Length 4419;
  Best Local Similarity
                         63.6%; Pred. No. 0.25;
 Matches
           49; Conservative
                                0; Mismatches
                                                 28;
                                                      Indels
                                                                    Gaps
                                                                            0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
                            | | | | | | | | | | | |
Db
        1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
           65 TTGTCACTTTCCGAGGA 81
Qу
             11 | 11 | 11 | 1 | 1 | 1
Db
        1605 TTCTCAGTTTTCCTGGA 1621
RESULT 24
US-10-161-539-3
; Sequence 3, Application US/10161539
; Publication No. US20030147854A1
   GENERAL INFORMATION:
        APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
        TITLE OF INVENTION: ADENOVIRUS VECTORS FOR GENE THERAPY
        NUMBER OF SEQUENCES: 10
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: GENZYME CORPORATION
             STREET: 15 PLEASANT STREET CONNECTOR
             CITY: FRAMINGHAM
             STATE: MASSACHUSETTS
             COUNTRY: USA
             ZIP: 01701-9322
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COMPUTER READABLE FORM:
             MEDIUM TYPE: Floppy disk
             COMPUTER: IBM PC compatible
             OPERATING SYSTEM: PC-DOS/MS-DOS
             SOFTWARE: ASCII
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/10/161,539
             FILING DATE: 20-Feb-2003
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: US 09/248,026
             FILING DATE: 10-FEB-1999
        ATTORNEY/AGENT INFORMATION:
             NAME: Newland, Bart G.
             REGISTRATION NUMBER: 31,282
             REFERENCE/DOCKET NUMBER: IG4-09.11.2-CON3
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: (508) 271-3920
             TELEFAX: (508) 872-5415
   INFORMATION FOR SEQ ID NO: 3:
        SEQUENCE CHARACTERISTICS:
             LENGTH: 5635 base pairs
             TYPE: nucleic acid
             STRANDEDNESS: single
             TOPOLOGY: linear
        MOLECULE TYPE: cDNA
        SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-10-161-539-3
 Query Match
                       31.6%; Score 32.2; DB 15; Length 5635;
 Best Local Similarity 63.6%; Pred. No. 0.27;
 Matches 49; Conservative 0; Mismatches 28; Indels
Qу
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
             Db
        2015 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 2074
          65 TTGTCACTTTCCGAGGA 81
Qу
             Db
        2075 TTCTCAGTTTTCCTGGA 2091
RESULT 25
US-09-982-315-3
; Sequence 3, Application US/09982315
; Publication No. US20030096762A1
; GENERAL INFORMATION:
; APPLICANT: Fischer, Horst
; APPLICANT: Illek, Beate
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
; FILE REFERENCE: 200116.403D1
 CURRENT APPLICATION NUMBER: US/09/982,315
  CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
  LENGTH: 6126
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TYPE: DNA
   ORGANISM: Homo sapiens
US-09-982-315-3
 Query Match
                       31.6%; Score 32.2; DB 10; Length 6126;
 Best Local Similarity 63.6%; Pred. No. 0.27;
 Matches 49; Conservative 0; Mismatches
                                            28;
                                                Indels
                                                             Gaps
                                                                    0;
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
         65 TTGTCACTTTCCGAGGA 81
Qу
            1605 TTCTCAGTTTTCCTGGA 1621
RESULT 26
US-09-782-378A-24
; Sequence 24, Application US/09782378A
; Patent No. US20020102731A1
; GENERAL INFORMATION:
  APPLICANT: Hearing, Patrick
  APPLICANT: Bahou, Wadie
  APPLICANT: Sandalon, Ziv
  APPLICANT: Gnatenko, Dmitri
  TITLE OF INVENTION: Adenoviral Vectors
; FILE REFERENCE: STONYB-04970
; CURRENT APPLICATION NUMBER: US/09/782,378A
; CURRENT FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/237,747
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 27
 SOFTWARE: PatentIn version 3.0
; SEQ ID NO 24
   LENGTH: 6129
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-782-378A-24
 Query Match
                       31.6%; Score 32.2; DB 9; Length 6129;
 Best Local Similarity 63.6%; Pred. No. 0.27;
 Matches 49; Conservative
                            0; Mismatches
                                            28; Indels
                                                         0; Gaps
                                                                    0;
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64 ,
Qy
            1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
         65 TTGTCACTTTCCGAGGA 81
Qу
            1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 27
US-09-982-315-1
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; Sequence 1, Application US/09982315

; Publication No. US20030096762A1

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; GENERAL INFORMATION:
  APPLICANT: Fischer, Horst
  APPLICANT: Illek, Beate
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
  FILE REFERENCE: 200116.403D1
  CURRENT APPLICATION NUMBER: US/09/982,315
  CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 6
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
   LENGTH: 6129
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-982-315-1
                       31.6%; Score 32.2; DB 10; Length 6129;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.27;
 Matches 49; Conservative 0; Mismatches 28; Indels
                                                          0; Gaps
                                                                    0;
          5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            1545 TATGGGAGAACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
         65 TTGTCACTTTCCGAGGA 81
Qy
            11 11 11 1 1 11
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 28
US-09-982-315-5
; Sequence 5, Application US/09982315
; Publication No. US20030096762A1
; GENERAL INFORMATION:
; APPLICANT: Fischer, Horst
  APPLICANT: Illek, Beate
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR CYSTIC FIBROSIS THERAPY
 FILE REFERENCE: 200116.403D1
; CURRENT APPLICATION NUMBER: US/09/982,315
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 6
 SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5
   LENGTH: 6129
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-982-315-5
                       31.6%; Score 32.2; DB 10; Length 6129;
 Query Match
 Best Local Similarity 63.6%; Pred. No. 0.27;
 Matches 49; Conservative 0; Mismatches 28; Indels
                                                           0; Gaps
                                                                     0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
            1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
Db
          65 TTGTCACTTTCCGAGGA 81
ΟV
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RESULT 29
US-10-161-539-1
; Sequence 1, Application US/10161539
 Publication No. US20030147854A1
   GENERAL INFORMATION:
        APPLICANT: Gregory, R.J., Armentano, D., Couture, L.A., Smith,
                   A.E.
        TITLE OF INVENTION: ADENOVIRUS VECTORS FOR GENE THERAPY
        NUMBER OF SEQUENCES: 10
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: GENZYME CORPORATION
             STREET: 15 PLEASANT STREET CONNECTOR
             CITY: FRAMINGHAM
             STATE: MASSACHUSETTS
             COUNTRY: USA
             ZIP: 01701-9322
        COMPUTER READABLE FORM:
             MEDIUM TYPE: Floppy disk
             COMPUTER: IBM PC compatible
             OPERATING SYSTEM: PC-DOS/MS-DOS
             SOFTWARE: ASCII
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/10/161,539
             FILING DATE: 20-Feb-2003
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: US 09/248,026
             FILING DATE: 10-FEB-1999
        ATTORNEY/AGENT INFORMATION:
             NAME: Newland, Bart G.
             REGISTRATION NUMBER: 31,282
             REFERENCE/DOCKET NUMBER: IG4-09.11.2-CON3
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: (508) 271-3920
             TELEFAX: (508) 872-5415
   INFORMATION FOR SEQ ID NO: 1:
        SEQUENCE CHARACTERISTICS:
             LENGTH: 6129 base pairs
             TYPE: nucleic acid
             STRANDEDNESS: single
             TOPOLOGY: linear
        MOLECULE TYPE: cDNA
        FEATURE:
             NAME/KEY: CDS
             LOCATION: 133..4572
        SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-161-539-1
                         31.6%; Score 32.2; DB 15; Length 6129;
 Query Match
                         63.6%;
                                Pred. No. 0.27;
  Best Local Similarity
                                                               0;
 Matches
           49; Conservative
                                0; Mismatches
                                                28;
                                                     Indels
                                                                   Gaps
                                                                           0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Οv
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Db
        1545 TATGGGAGACTGGAGCCTTCAGAGGGTAAAATTAAGCACAGTGGAAGAATTTCATTCTG 1604
          65 TTGTCACTTTCCGAGGA 81
QУ
             11 111 111 1 111
        1605 TTCTCAGTTTTCCTGGA 1621
Db
RESULT 30
US-10-369-493-37753/c
; Sequence 37753, Application US/10369493
; Publication No. US20030233675A1
; GENERAL INFORMATION:
; APPLICANT: Cao, Yongwei
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Goldman, Barry S.
  APPLICANT: Chen, Xianfeng
  TITLE OF INVENTION: EXPRESSION OF MICROBIAL PROTEINS IN PLANTS FOR PRODUCTION
OF
  TITLE OF INVENTION: PLANTS WITH IMPROVED PROPERTIES
  FILE REFERENCE: 38-10(52052)B
  CURRENT APPLICATION NUMBER: US/10/369,493
  CURRENT FILING DATE: 2003-02-28
  PRIOR APPLICATION NUMBER: US 60/360,039
 PRIOR FILING DATE: 2002-02-21
  NUMBER OF SEQ ID NOS: 47374
; SEQ ID NO 37753
   LENGTH: 2310
   TYPE: DNA
   ORGANISM: Pseudomonas fluorescens
US-10-369-493-37753
                        28.4%; Score 29; DB 16; Length 2310;
 Query Match
 Best Local Similarity 57.0%; Pred. No. 3.1;
                            0; Mismatches 40; Indels
 Matches 53; Conservative
                                                             0; Gaps
                                                                         0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qy
             374 TTGGTCAGCTCTTGGGCGTATTGAGTCTTGCCGTGCTCACCGCCGCAGCAGGACGACGTG 315
Db
         65 TTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
QУ
               1 111
                     314 GCGCCACGGGCCATGGCAAACAAGGTGTCGAGG 282
Dh
RESULT 31
US-10-027-632-172129
; Sequence 172129, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
  CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
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PRIOR FILING DATE: 2000-07-12
   PRIOR APPLICATION NUMBER: US 60/198,676
   PRIOR FILING DATE: 2000-04-20
   PRIOR APPLICATION NUMBER: US 60/193,483
   PRIOR FILING DATE: 2000-03-29
   PRIOR APPLICATION NUMBER: US 60/185,218
   PRIOR FILING DATE: 2000-02-24
   PRIOR APPLICATION NUMBER: US 60/167,363
   PRIOR FILING DATE: 1999-11-23
   PRIOR APPLICATION NUMBER: US 60/156,358
   PRIOR FILING DATE: 1999-09-28
   PRIOR APPLICATION NUMBER: US 60/146,002
   PRIOR FILING DATE: 1999-08-09
   NUMBER OF SEQ ID NOS: 325720
   SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 172129
    LENGTH: 799
    TYPE: DNA
    ORGANISM: Human
US-10-027-632-172129
  Query Match
                         27.8%;
                                 Score 28.4; DB 13; Length 799;
  Best Local Similarity
                         62.9%; Pred. No. 3.9;
                                0; Mismatches
           44; Conservative
                                                 26;
                                                      Indels
                                                                0;
                                                                   Gaps
                                                                            0;
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
Qу
              Db
         591 TAGGTAATATCAGTGTGCTCCAAAGGTTGAGAATAACTGCTTTAAGTTGAAAAAAAGAATG 650
          65 TTGTCACTTT 74
Qу
              Db
         651 TTGGAACTCT 660
RESULT 32
US-10-027-632-172129
; Sequence 172129, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
  CURRENT APPLICATION NUMBER: US/10/027,632
   CURRENT FILING DATE: 2002-04-30
   PRIOR APPLICATION NUMBER: US 60/218,006
   PRIOR FILING DATE: 2000-07-12
   PRIOR APPLICATION NUMBER: US 60/198,676
   PRIOR FILING DATE: 2000-04-20
   PRIOR APPLICATION NUMBER: US 60/193,483
   PRIOR FILING DATE: 2000-03-29
   PRIOR APPLICATION NUMBER: US 60/185,218
   PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
   PRIOR FILING DATE: 1999-11-23
   PRIOR APPLICATION NUMBER: US 60/156,358
   PRIOR FILING DATE: 1999-09-28
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; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 172129
   LENGTH: 799
   TYPE: DNA
   ORGANISM: Human
US-10-027-632-172129
 Query Match 27.8%; Score 28.4; DB 16; Length 799; Best Local Similarity 62.9%; Pred. No. 3.9;
 Matches 44; Conservative 0; Mismatches
                                                26; Indels
                                                               0; Gaps
                                                                           0:
Qy
           5 TAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAGACTG 64
             591 TAGGTAATATCAGTGTGCTCCAAAGGTTGAGAATAACTGCTTTAAGTTGAAAAAAAGAATG 650
          65 TTGTCACTTT 74
Qу
             Db
         651 TTGGAACTCT 660
RESULT 33
US-09-925-299-368/c
; Sequence 368, Application US/09925299
; Patent No. US20020055627A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
 TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA102
; CURRENT APPLICATION NUMBER: US/09/925,299
  CURRENT FILING DATE: 2001-08-10
  PRIOR APPLICATION NUMBER: PCT/US00/05883
  PRIOR FILING DATE: 2000-03-08
  PRIOR APPLICATION NUMBER: 60/124,270
  PRIOR FILING DATE: 1999-03-12
  NUMBER OF SEQ ID NOS: 1556
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 368
   LENGTH: 548
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (370)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (378)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (384)
   OTHER INFORMATION: n equals a,t,q, or c
   NAME/KEY: misc feature
   LOCATION: (412)
   OTHER INFORMATION: n equals a,t,q, or c
   NAME/KEY: misc feature
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LOCATION: (429)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (449)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (471)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (490)
   OTHER INFORMATION: n equals a,t,q, or c
   NAME/KEY: misc feature
   LOCATION: (495)
   OTHER INFORMATION: n equals a,t,q, or c
   NAME/KEY: misc feature
   LOCATION: (528)
   OTHER INFORMATION: n equals a,t,g, or c
US-09-925-299-368
 Query Match
                         27.6%; Score 28.2; DB 9; Length 548;
 Best Local Similarity 61.6%; Pred. No. 4.2;
 Matches 45; Conservative
                               0; Mismatches
                                                28; Indels
                                                               0; Gaps
                                                                          0;
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
Qу
             327 GTGTTGGTGTGACTATTGTAGCTGGGACATTTACTGTGGTGGGTTTCTGAGGAGTTGGTG 268
Db
          89 CTGTCCTGGAGGC 101
Qу
               267 GGGTTCTTGTAGC 255
Db
RESULT 34
US-09-925-299-368/c
; Sequence 368, Application US/09925299
; Publication No. US20030040617A9
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
  TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
  FILE REFERENCE: PA102
  CURRENT APPLICATION NUMBER: US/09/925,299
  CURRENT FILING DATE: 2001-08-10
  PRIOR APPLICATION NUMBER: PCT/US00/05883
  PRIOR FILING DATE: 2000-03-08
  PRIOR APPLICATION NUMBER: 60/124,270
  PRIOR FILING DATE: 1999-03-12
  NUMBER OF SEQ ID NOS: 1556
  SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO 368
   LENGTH: 548
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (370)
   OTHER INFORMATION: n equals a,t,q, or c
   NAME/KEY: misc feature
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LOCATION: (378)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (384)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (412)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (429)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (449)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (471)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (490)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (495)
   OTHER INFORMATION: n equals a,t,g, or c
   NAME/KEY: misc feature
   LOCATION: (528)
   OTHER INFORMATION: n equals a,t,q, or c
US-09-925-299-368
                        27.6%; Score 28.2; DB 10; Length 548;
  Query Match
                        61.6%; Pred. No. 4.2;
  Best Local Similarity
                                                              0; Gaps
                                                                         0;
                               0; Mismatches
                                              28; Indels
 Matches 45; Conservative
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
Qу
             327 GTGTTGGTGTGACTATTGTAGCTGGGACATTTACTGTGGTGGGTTTCTGAGGAGTTGGTG 268
          89 CTGTCCTGGAGGC 101
Qу
               Db
         267 GGGTTCTTGTAGC 255
RESULT 35
US-09-934-814-4/c
; Sequence 4, Application US/09934814
; Patent No. US20020137159A1
; GENERAL INFORMATION:
; APPLICANT: Lok, Si
; APPLICANT: Holloway, James L.
 APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/09/934,814
; CURRENT FILING DATE: 2001-08-22
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: FastSEO for Windows Version 3.0
; SEQ ID NO 4
   LENGTH: 432
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TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(429)
US-09-934-814-4
                        27.5%; Score 28; DB 9; Length 432;
 Query Match
                        71.2%; Pred. No. 4.7;
 Best Local Similarity
                                               15; Indels
                                                             0; Gaps
 Matches
          37; Conservative
                               0; Mismatches
          46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
             366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
Db
RESULT 36
US-10-142-465-4/c
; Sequence 4, Application US/10142465
; Publication No. US20030166070A1
; GENERAL INFORMATION:
  APPLICANT: Lok, Si
 APPLICANT: Holloway, James L.
  APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/10/142,465
  CURRENT FILING DATE: 2002-05-09
  NUMBER OF SEQ ID NOS: 13
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEO ID NO 4
   LENGTH: 432
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(429)
US-10-142-465-4
  Query Match
                        27.5%; Score 28; DB 15; Length 432;
                        71.2%; Pred. No. 4.7;
  Best Local Similarity
          37; Conservative
                               0; Mismatches
                                               15; Indels
                                                             0;
                                                                 Gaps
                                                                        0;
 Matches
          46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
             111 1111
         366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGCCACTGTGTTCTGG 315
Db
RESULT 37
US-09-934-814-1/c
; Sequence 1, Application US/09934814
; Patent No. US20020137159A1
; GENERAL INFORMATION:
; APPLICANT: Lok, Si
 APPLICANT: Holloway, James L.
 APPLICANT: O'Hara, Patrick J.
; TITLE OF INVENTION: Human Phermone Polypepides
```

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FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/09/934,814
  CURRENT FILING DATE: 2001-08-22
; NUMBER OF SEQ ID NOS: 13
 SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 1
   LENGTH: 525
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(525)
US-09-934-814-1
 Query Match
                       27.5%; Score 28; DB 9; Length 525;
 Best Local Similarity 71.2%; Pred. No. 4.9;
 Matches 37; Conservative 0; Mismatches 15; Indels 0; Gaps
         46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
            Db
         375 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 324
RESULT 38
US-10-142-465-1/c
; Sequence 1, Application US/10142465
; Publication No. US20030166070A1
; GENERAL INFORMATION:
; APPLICANT: Lok, Si
; APPLICANT: Holloway, James L.
; APPLICANT: O'Hara, Patrick J.
; TITLE OF INVENTION: Human Phermone Polypepides
; FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/10/142,465
  CURRENT FILING DATE: 2002-05-09
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 1
   LENGTH: 525
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(525)
US-10-142-465-1
                       27.5%; Score 28; DB 15; Length 525;
 Query Match
 Best Local Similarity 71.2%; Pred. No. 4.9;
 Matches 37; Conservative 0; Mismatches 15; Indels
                                                          0; Gaps 0;
         46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qy
            375 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 324
RESULT 39
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RESULT 39 US-09-934-814-7/c

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; Sequence 7, Application US/09934814
; Patent No. US20020137159A1
; GENERAL INFORMATION:
  APPLICANT: Lok, Si
  APPLICANT: Holloway, James L.
  APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/09/934,814
  CURRENT FILING DATE: 2001-08-22
  NUMBER OF SEQ ID NOS: 13
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 7
   LENGTH: 540
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(537)
US-09-934-814-7
 Query Match
                         27.5%;
                                Score 28; DB 9; Length 540;
 Best Local Similarity
                        71.2%; Pred. No. 5;
                                                               0; Gaps
                               0; Mismatches
                                                                          0;
          37; Conservative
                                                15; Indels
          46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
             Db
         366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
RESULT 40
US-10-142-465-7/c
; Sequence 7, Application US/10142465
; Publication No. US20030166070A1
; GENERAL INFORMATION:
  APPLICANT: Lok, Si
  APPLICANT: Holloway, James L.
 APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/10/142,465
  CURRENT FILING DATE: 2002-05-09
  NUMBER OF SEQ ID NOS: 13
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 7
   LENGTH: 540
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(537)
US-10-142-465-7
 Query Match
                         27.5%; Score 28; DB 15; Length 540;
 Best Local Similarity 71.2%; Pred. No. 5;
 Matches 37; Conservative
                               0; Mismatches
                                                15; Indels
                                                               0; Gaps
                                                                          0;
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```
46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
             366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
Db
RESULT 41
US-09-934-814-10/c
; Sequence 10, Application US/09934814
; Patent No. US20020137159A1
; GENERAL INFORMATION:
; APPLICANT: Lok, Si
; APPLICANT: Holloway, James L.
; APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/09/934,814
  CURRENT FILING DATE: 2001-08-22
  NUMBER OF SEQ ID NOS: 13
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 10
   LENGTH: 795
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1)...(792)
US-09-934-814-10
                        27.5%; Score 28; DB 9; Length 795;
 Query Match
 Best Local Similarity 71.2%; Pred. No. 5.5;
 Matches 37; Conservative
                             0; Mismatches 15; Indels
                                                             0; Gaps
                                                                        0;
          46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
Qу
             366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
Db
RESULT 42
US-10-142-465-10/c
; Sequence 10, Application US/10142465
; Publication No. US20030166070A1
; GENERAL INFORMATION:
; APPLICANT: Lok, Si
  APPLICANT: Holloway, James L.
  APPLICANT: O'Hara, Patrick J.
  TITLE OF INVENTION: Human Phermone Polypepides
  FILE REFERENCE: 00-80
  CURRENT APPLICATION NUMBER: US/10/142,465
  CURRENT FILING DATE: 2002-05-09
  NUMBER OF SEQ ID NOS: 13
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 10
   LENGTH: 795
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
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; LOCATION: (1)...(792)
US-10-142-465-10
                        27.5%; Score 28; DB 15; Length 795;
  Query Match
 Best Local Similarity 71.2%; Pred. No. 5.5;
 Matches 37; Conservative 0; Mismatches 15; Indels
                                                            0; Gaps
                                                                     0;
Qу
         46 GTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAGCTGTCCTGG 97
             366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
Db
RESULT 43
US-10-399-456-3/c
; Sequence 3, Application US/10399456
; Publication No. US20040043395A1
; GENERAL INFORMATION:
; APPLICANT: INCYTE GENOMICS, INC.
; APPLICANT: LAL, Preeti G.
; APPLICANT: CHAWLA, Narinder K.
; APPLICANT: GANDHI, Ameena R.
; APPLICANT: LU, Yan
; APPLICANT: RAMKUMAR, Jayalaxmi
; APPLICANT: BAUGHN, Mariah R.
 APPLICANT: BRUNS, Christopher M.
 APPLICANT: HAFALIA, April J.A.
; APPLICANT: YAO, Monique G.
  TITLE OF INVENTION: LIPOCALINS
; FILE REFERENCE: PF-0822 USN
; CURRENT APPLICATION NUMBER: US/10/399,456
; CURRENT FILING DATE: 2003-04-14
; PRIOR APPLICATION NUMBER: PCT/US01/31942
; PRIOR FILING DATE: 2001-10-12
; PRIOR APPLICATION NUMBER: US 60/240,541
; PRIOR FILING DATE: 2000-10-13
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: PERL Program
; SEO ID NO 3
  LENGTH: 1630
  TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   OTHER INFORMATION: Incyte ID No. US20040043395A1 3537562CB1
US-10-399-456-3
 Query Match
                        27.5%; Score 28; DB 13; Length 1630;
 Best Local Similarity 71.2%; Pred. No. 6.6;
 Matches 37; Conservative 0; Mismatches 15; Indels
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Qу
             366 GAAGGTGATGAACAGCCTGTAGTCAGTCTCCGAGACGGCCACTGTGTTCTGG 315
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RESULT 44 US-10-052-482-166

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; Sequence 166, Application US/10052482
; Publication No. US20040072264A1
; GENERAL INFORMATION:
  APPLICANT: Engelhard, Eric
  APPLICANT: Morris, David
  TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
  FILE REFERENCE: A-71087/RMS/DCF
  CURRENT APPLICATION NUMBER: US/10/052,482
  CURRENT FILING DATE: 2002-08-15
  PRIOR APPLICATION NUMBER: US 09/747,377
  PRIOR FILING DATE: 2000-12-22
  PRIOR APPLICATION NUMBER: US 09/798,586
  PRIOR FILING DATE: 2001-03-02
  NUMBER OF SEQ ID NOS: 241
  SOFTWARE: PatentIn version 3.1
; SEQ ID NO 166
   LENGTH: 48244
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc_feature
   LOCATION: (36673)..(36711)
   OTHER INFORMATION: "n" at positions 36673 to 36711 can be any base
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (40035)..(40119)
   OTHER INFORMATION: "n" at positions 40035 to 40119 can be any base
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (42958)..(43306)
   OTHER INFORMATION: "n" at positions 42958 to 43306 can be any base
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (47841)..(47909)
   OTHER INFORMATION: "n" at positions 47841 to 47909 can be any base
US-10-052-482-166
                         27.5%; Score 28; DB 12; Length 48244;
 Query Match
 Best Local Similarity 58.3%; Pred. No. 16;
           49; Conservative
                               0; Mismatches
                                                35; Indels
                                                                          0;
           1 CTGGTAGGTGAGATCTCTGACCTCCAGAGTGTTGGACTGACCACTGTAGGTGAAGTACAG 60
Qу
                   Db
       40187 CTGGCTTGTGTGCTTCCTGCCCTCCACTGGGTGCTACGGACCAAGGGCTGTGCTGAGCCC 40246
          61 ACTGTTGTCACTTTCCGAGGAGAA 84
Qу
              Db
       40247 CCTGTGGCCGCTCTCACAGCTGAA 40270
RESULT 45
US-10-027-632-265948
; Sequence 265948, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
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TITLE OF INVENTION: Polymorphisms in the Human Genome
   FILE REFERENCE: 108827.129
   CURRENT APPLICATION NUMBER: US/10/027,632
   CURRENT FILING DATE: 2002-04-30
   PRIOR APPLICATION NUMBER: US 60/218,006
   PRIOR FILING DATE: 2000-07-12
   PRIOR APPLICATION NUMBER: US 60/198,676
   PRIOR FILING DATE: 2000-04-20
   PRIOR APPLICATION NUMBER: US 60/193,483
   PRIOR FILING DATE: 2000-03-29
   PRIOR APPLICATION NUMBER: US 60/185,218
   PRIOR FILING DATE: 2000-02-24
   PRIOR APPLICATION NUMBER: US 60/167,363
   PRIOR FILING DATE: 1999-11-23
   PRIOR APPLICATION NUMBER: US 60/156,358
   PRIOR FILING DATE: 1999-09-28
   PRIOR APPLICATION NUMBER: US 60/146,002
   PRIOR FILING DATE: 1999-08-09
   NUMBER OF SEQ ID NOS: 325720
   SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 265948
   LENGTH: 987
   TYPE: DNA
   ORGANISM: Human
    FEATURE:
    NAME/KEY: misc feature
    LOCATION: (1)...(987)
    OTHER INFORMATION: n = A, T, C or G
US-10-027-632-265948
  Query Match
                         27.3%; Score 27.8; DB 13; Length 987;
  Best Local Similarity
                         62.0%; Pred. No. 6.9;
           44; Conservative
                                0; Mismatches
                                                 27; Indels
                                                                0; Gaps
                                                                           0;
Qу
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
                      Db
         253 GTGGTGGTAGGAGCATTGCAGAGGATGGACAGATTCATGTCCCTCAGAGAGGAGGAGGAGGAG 312
          89 CTGTCCTGGAG 99
Qу
               1 1 1 111
Db
          313 AAGGCATAGAG 323
RESULT 46
US-10-027-632-265949
; Sequence 265949, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
   FILE REFERENCE: 108827.129
   CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
  PRIOR APPLICATION NUMBER: US 60/218,006
  PRIOR FILING DATE: 2000-07-12
  PRIOR APPLICATION NUMBER: US 60/198,676
```

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PRIOR FILING DATE: 2000-04-20
  PRIOR APPLICATION NUMBER: US 60/193,483
  PRIOR FILING DATE: 2000-03-29
  PRIOR APPLICATION NUMBER: US 60/185,218
  PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
  PRIOR FILING DATE: 1999-11-23
  PRIOR APPLICATION NUMBER: US 60/156,358
  PRIOR FILING DATE: 1999-09-28
  PRIOR APPLICATION NUMBER: US 60/146,002
  PRIOR FILING DATE: 1999-08-09
  NUMBER OF SEQ ID NOS: 325720
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 265949
   LENGTH: 987
   TYPE: DNA
   ORGANISM: Human
   FEATURE:
   NAME/KEY: misc_feature
   LOCATION: (1)...(987)
   OTHER INFORMATION: n = A, T, C or G
US-10-027-632-265949
  Query Match
                        27.3%; Score 27.8; DB 13; Length 987;
 Best Local Similarity
                        62.0%; Pred. No. 6.9;
 Matches 44; Conservative
                              0; Mismatches
                                              27;
                                                   Indels
                                                            0; Gaps
                                                                        0; .
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
Qy
             Db
         89 CTGTCCTGGAG 99
Qv
               Db
         313 AAGGCATAGAG 323
RESULT 47
US-10-027-632-265950
; Sequence 265950, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
  CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
  PRIOR APPLICATION NUMBER: US 60/218,006
  PRIOR FILING DATE: 2000-07-12
  PRIOR APPLICATION NUMBER: US 60/198,676
  PRIOR FILING DATE: 2000-04-20
  PRIOR APPLICATION NUMBER: US 60/193,483
  PRIOR FILING DATE: 2000-03-29
  PRIOR APPLICATION NUMBER: US 60/185,218
  PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
  PRIOR FILING DATE: 1999-11-23
```

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PRIOR APPLICATION NUMBER: US 60/156,358
   PRIOR FILING DATE: 1999-09-28
   PRIOR APPLICATION NUMBER: US 60/146,002
   PRIOR FILING DATE: 1999-08-09
   NUMBER OF SEQ ID NOS: 325720
   SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 265950
   LENGTH: 987
    TYPE: DNA
    ORGANISM: Human
    FEATURE:
    NAME/KEY: misc feature
    LOCATION: (1)...(987)
    OTHER INFORMATION: n = A, T, C or G
US-10-027-632-265950
  Query Match
                         27.3%; Score 27.8; DB 13; Length 987;
  Best Local Similarity 62.0%; Pred. No. 6.9;
  Matches
           44; Conservative
                                0; Mismatches
                                                 27;
                                                      Indels
Qγ
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
                      Db
         253 GTGGTGGTAGGAGCATTGCAGAGGATGGACAGATTCATGTCCCTCAGAGAGGAGGAGGAG 312
          89 CTGTCCTGGAG 99
Qу
               1 1 1 1 1
Db
         313 AAGGCATAGAG 323
RESULT 48
US-10-027-632-265951
; Sequence 265951, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
   CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
  PRIOR APPLICATION NUMBER: US 60/218,006
  PRIOR FILING DATE: 2000-07-12
  PRIOR APPLICATION NUMBER: US 60/198,676
  PRIOR FILING DATE: 2000-04-20
  PRIOR APPLICATION NUMBER: US 60/193,483
  PRIOR FILING DATE: 2000-03-29
  PRIOR APPLICATION NUMBER: US 60/185,218
  PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
  PRIOR FILING DATE: 1999-11-23
  PRIOR APPLICATION NUMBER: US 60/156,358
  PRIOR FILING DATE: 1999-09-28
  PRIOR APPLICATION NUMBER: US 60/146,002
  PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 265951
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LENGTH: 987
    TYPE: DNA
    ORGANISM: Human
    FEATURE:
    NAME/KEY: misc feature
    LOCATION: (1)...(987)
    OTHER INFORMATION: n = A, T, C or G
US-10-027-632-265951
  Query Match
                         27.3%; Score 27.8; DB 13; Length 987;
  Best Local Similarity
                         62.0%; Pred. No. 6.9;
  Matches 44; Conservative
                                0; Mismatches
                                                 27; Indels
                                                                0;
                                                                    Gaps
                                                                            0;
           29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
Qу
                      253 GTGGTGGTAGGAGCATTGCAGAGGATGGACAGATTCATGTCCCTCAGAGAGGAGGAGGAG
Db
Qу
          89 CTGTCCTGGAG 99
                1 | | | | |
Db
          313 AAGGCATAGAG 323
RESULT 49
US-10-027-632-265948
; Sequence 265948, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
  CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
  PRIOR APPLICATION NUMBER: US 60/218,006
  PRIOR FILING DATE: 2000-07-12
  PRIOR APPLICATION NUMBER: US 60/198,676
  PRIOR FILING DATE: 2000-04-20
  PRIOR APPLICATION NUMBER: US 60/193,483
  PRIOR FILING DATE: 2000-03-29
  PRIOR APPLICATION NUMBER: US 60/185,218
  PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
  PRIOR FILING DATE: 1999-11-23
  PRIOR APPLICATION NUMBER: US 60/156,358
  PRIOR FILING DATE: 1999-09-28
  PRIOR APPLICATION NUMBER: US 60/146,002
  PRIOR FILING DATE: 1999-08-09
  NUMBER OF SEQ ID NOS: 325720
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 265948
   LENGTH: 987
   TYPE: DNA
   ORGANISM: Human
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (1)...(987)
   OTHER INFORMATION: n = A, T, C or G
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Qу

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Query Match
                       27.3%; Score 27.8; DB 16; Length 987;
  Best Local Similarity 62.0%; Pred. No. 6.9;
  Matches 44; Conservative
                            0; Mismatches
                                              27; Indels
                                                            0; Gaps
                                                                       0;
          29 GTGTTGGACTGACCACTGTAGGTGAAGTACAGACTGTTGTCACTTTCCGAGGAGAACAAG 88
Qy
                     Db
         89 CTGTCCTGGAG 99
Qγ
               1 1 1 111
         313 AAGGCATAGAG 323
Db
RESULT 50
US-10-027-632-265949
; Sequence 265949, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
  TITLE OF INVENTION: Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.129
  CURRENT APPLICATION NUMBER: US/10/027,632
  CURRENT FILING DATE: 2002-04-30
  PRIOR APPLICATION NUMBER: US 60/218,006
  PRIOR FILING DATE: 2000-07-12
  PRIOR APPLICATION NUMBER: US 60/198,676
  PRIOR FILING DATE: 2000-04-20
  PRIOR APPLICATION NUMBER: US 60/193,483
  PRIOR FILING DATE: 2000-03-29
  PRIOR APPLICATION NUMBER: US 60/185,218
  PRIOR FILING DATE: 2000-02-24
  PRIOR APPLICATION NUMBER: US 60/167,363
  PRIOR FILING DATE: 1999-11-23
  PRIOR APPLICATION NUMBER: US 60/156,358
  PRIOR FILING DATE: 1999-09-28
  PRIOR APPLICATION NUMBER: US 60/146,002
  PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEO ID NO 265949
   LENGTH: 987
   TYPE: DNA
   ORGANISM: Human
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (1)...(987)
   OTHER INFORMATION: n = A, T, C or G
US-10-027-632-265949
 Query Match
                       27.3%; Score 27.8; DB 16; Length 987;
 Best Local Similarity 62.0%; Pred. No. 6.9;
 Matches 44; Conservative
                              0; Mismatches
                                             27; Indels
                                                               Gaps
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Search completed: April 29, 2004, 21:08:43 Job time: 100.194 secs